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(71) Applicant (for all designated States except US): HUMAN GENOME SCIENCES, INC. [US/US]; 9410 Key West Avenue, Rockville, MD 20850 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): YOUNG, Paul [US/US];
122 Beckwith Street, Gaithersburg, MD 20878 (US).
GREENE, John, M. [US/US]; 872 Diamond Drive,
Gaithersburg, MD 20878 (US). FERRIE, Ann, M. [US/US];
13203 L Astoria Hill Court, Germantown, MD 20874 (US).
RUBEN, Steven, M. [US/US]; 18528 Heritage Hills Drive,
Olney, MD 20832 (US). ROSEN, Craig, A. [US/US];
22400 Rolling Hill Road, Laytonsville, MD 20882 (US).
HU, Jing-Shan [CN/US]; 1247 Lakeside Drive #3034,
Sunnyvale, CA 94086 (US). OLSEN, Henrik, S. [DK/US];
182 Kendrick Place #24, Gaithersburg, MD 20878 (US).
EBNER, Reinhard [DE/US]; 9906 Shelburne Terrace #316,

Gaithersburg, MD 20878 (US). BREWER, Laurie, A. [US/US]; 14920 Mt. Nebo Road, Poolesville, MD 20837 (US). MOORE, Paul, A. [GB/US]; Apartment 104, 1908 Holly Ridge Drive, McLean, VA 22102 (US). SHI, Yanggu [CN/US]; 437 West Side Drive, Gaithersburg, MD 20878 (US). FLORENCE, Charles [US/US]; (US). FLORENCE, Kimberly [US/US]; 12805 Atlantic Avenue, Rockville, MD 20851 (US). LAFLEUR, David, W. [US/US]; 1615 Q Street, N.W. #807, Washington, DC 20009 (US). NI, Jian [CN/US]; 5502 Manorfield Road, Rockville, MD 20853 (US). FAN, Ping [CN/US]; Apartment 302, 335 West Side Drive, Gaithesburg, MD 20878 (US). WEI, Ying-Fei [CN/US]; 13524 Straw Bale Lane, Darnestown, MD 20878 (US). FISCHER, Carrie, L. [US/US]; 5810 Hall Street, Burke, VA 22015 (US). SOPPET, Daniel, R. [US/US]; 15050 Stillfield Place, Centreville, VA 22020 (US). LI, Yi [CN/US]; 1247 Lakeside Drive #3034, Sunnyvale, CA 94086 (US). ZENG, Zhizhen [CN/US]; 13950 Saddleview Drive, Gaithersburg, MD 20878 (US). KYAW, Hla [MM/US]; 520 Sugarbush Circle, Frederick, MD 21703 (US). YU, Guo-Liang [CN/US]; 13524 Straw Bale Lane, Darnestown, MD 20878 (US). FENG, Ping [CN/US]; 4 Relda Court, Gaithersburg, MD 20878 (US). DILLON, Patrick, J. [US/US]; 1055 Snipe Court, Carlsbad, CA 92009 (US). ENDRESS, Gregory, A. [US/US]; 9729 Clagett Farm Drive, Potomac, MD 20854 (US). CARTER, Kenneth, C. [US/US]; 11601 Brandy Hall Lane, North Potomac, MD 20878 (US).

- (74) Agents: HOOVER, Kenley, K. et al.; Human Genome Sciences, Inc., 9410 Key West Avenue, Rockville, MD 10850 (US).
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# **207 Human Secreted Proteins**

# Field of the Invention

This invention relates to newly identified polynucleotides and the polypeptides encoded by these polynucleotides, uses of such polynucleotides and polypeptides, and their production.

# Background of the Invention

Unlike bacterium, which exist as a single compartment surrounded by a membrane, human cells and other eucaryotes are subdivided by membranes into many functionally distinct compartments. Each membrane-bounded compartment, or organelle, contains different proteins essential for the function of the organelle. The cell uses "sorting signals," which are amino acid motifs located within the protein, to target proteins to particular cellular organelles.

One type of sorting signal, called a signal sequence, a signal peptide, or a leader sequence, directs a class of proteins to an organelle called the endoplasmic reticulum (ER). The ER separates the membrane-bounded proteins from all other types of proteins. Once localized to the ER, both groups of proteins can be further directed to another organelle called the Golgi apparatus. Here, the Golgi distributes the proteins to vesicles, including secretory vesicles, the cell membrane, lysosomes, and the other organelles.

Proteins targeted to the ER by a signal sequence can be released into the extracellular space as a secreted protein. For example, vesicles containing secreted proteins can fuse with the cell membrane and release their contents into the extracellular space - a process called exocytosis. Exocytosis can occur constitutively or after receipt of a triggering signal. In the latter case, the proteins are stored in secretory vesicles (or secretory granules) until exocytosis is triggered. Similarly, proteins residing on the cell membrane can also be secreted into the extracellular space by proteolytic cleavage of a "linker" holding the protein to the membrane.

Despite the great progress made in recent years, only a small number of genes encoding human secreted proteins have been identified. These secreted proteins include the commercially valuable human insulin, interferon, Factor VIII, human growth hormone, tissue plasminogen activator, and erythropoeitin. Thus, in light of the pervasive role of secreted proteins in human physiology, a need exists for identifying and characterizing novel human secreted proteins and the genes that encode them. This knowledge will allow one to detect, to treat, and to prevent medical disorders by using secreted proteins or the genes that encode them.

### Summary of the Invention

The present invention relates to novel polynucleotides and the encoded polypeptides. Moreover, the present invention relates to vectors, host cells, antibodies, and recombinant methods for producing the polypeptides and polynucleotides. Also provided are diagnostic methods for detecting disorders related to the polypeptides, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying binding partners of the polypeptides.

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# Detailed Description

#### **Definitions**

The following definitions are provided to facilitate understanding of certain terms used throughout this specification.

In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide.

In the present invention, a "secreted" protein refers to those proteins capable of being directed to the ER, secretory vesicles, or the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily containing a signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage.

As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence contained in SEQ ID NO:X or the cDNA contained within the clone deposited with the ATCC. For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, with or without the signal sequence, the secreted protein coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having the translated amino acid sequence generated from the polynucleotide as broadly defined.

In the present invention, the full length sequence identified as SEQ ID NO:X was often generated by overlapping sequences contained in multiple clones (contig

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analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X was deposited with the American Type Culture Collection ("ATCC"). As shown in Table 1, each clone is identified by a cDNA Clone ID (Identifier) and the ATCC Deposit Number. The ATCC is located at 10801 University Boulevard, Manassas, Virginia 20110-2209, USA. The ATCC deposit was made pursuant to the terms of the Budapest Treaty on the international recognition of the deposit of microorganisms for purposes of patent procedure.

A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, the complement thereof, or the cDNA within the clone deposited with the ATCC. "Stringent hybridization conditions" refers to an overnight incubation at 42° C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 μg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH<sub>2</sub>PO<sub>4</sub>; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a

WO 98/54963

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complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine. A variety of modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.

The polypeptide of the present invention can be composed of amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphotidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine,

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formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth Enzymol 182:626-646 (1990); Rattan et al., Ann NY Acad Sci 663:48-62 (1992).)

"SEQ ID NO:X" refers to a polynucleotide sequence while "SEQ ID NO:Y" refers to a polypeptide sequence, both sequences identified by an integer specified in Table 1.

"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.)

# Polynucleotides and Polypeptides of the Invention

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 1

This gene is expressed primarily in melanocytes and, to a lesser extent, in testes, ovary, kidney and other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer, disorders of neural crest derived cells including pigmentation defects, melanoma, reproductive organ defects, and defects of the kidney. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skin,

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reproductive, and renal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating disorders that arise from alterations in the number or fate of neural crest derived cells including cancers such as melanoma and defects of the developing reproductive system.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 2

This gene is expressed primarily in infant brain and fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental disorders of the brain or lung. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous and pulmonary systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating or diagnosing disorders associated with abnormal proliferation of cells in the Central nervous system and developing lung.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 3

This gene is expressed primarily in breast lymph node and to a lesser extent in ovarian cancer and chondrosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune responses such as inflammation or immune surveillance for

tumors. This gene may be important for inflammatory responses associated with tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 236 as residues: Lys-45 to Val-50, Lys-69 to Arg-76.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment or diagnosis of immune responses including those associated with tumor-induced inflammation.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 4

This gene is expressed primarily in T-cells and T-cell lymphomas.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunilogical diseases involving T-cells such as inflammation, autoimmunity, and cancers including T-cell lymphomas. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of T-cells and other cells of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing and treating T-cell based disorders such as inflammatory diseases, autoimmune disease and tumors including T-cell lymphomas.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 5

This gene is expressed primarily in activated monocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation, autoimmunity, infection, or disorders involving activation of monocytes. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 238 as residues: Asp-19 to Arg-31.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing or treating diseases that result in activation of monocytes including infections, inflammatory responses or autoimmune diseases.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 6

The translation product of this gene shares sequence homology with terminal deoxynucleotidyltransferase which is thought to be important in catalyzing the elongation of oligo- or polydeoxynucleotide chains.

This gene is expressed primarily in activated human neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer, particularly those of the blood such as leukemia and deficiencies in neutrophils such as neutropenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having

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such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to terminal deoxynucleotidyltransferase indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and differential diagnosis of acute leukemia's. Alternatively, this gene may function in the proliferation of neutrophils and be useful as a treatment for neutropenia, for example, following neutropenia as a result of chemotherapy.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 7

The contig exhibits a reasonable homology to the human chorionic gonadotropic (HCG) analogue-GT beta-subunit as disclosed in U.S. Patent No. 5,508,261 and PCT Publication No. WO 92/22568. There is a high degree of conservation of the structurally important cysteine residues in these identities.

This gene is expressed primarily in IL-1 and LPS induced neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system, including inflammatory diseases and allergies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of diseases of the immune system since expression is primarily in neutrophils, and may be useful as a growth factor for the differentiation or proliferation of neutrophils for the treatment of neutropenia following chemotherapy.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 8

This gene is expressed primarily in IL-1- and LPS-induced neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system, including inflammatory diseases and allergies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 241 as residues: Ser-14 to Pro-22, Leu-43 to Val-53.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of diseases of the immune system since expression is primarily in neutrophils, and may be useful as a growth factor for the differentiation or proliferation of neutrophils for the treatment of neutropenia following chemotherapy.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 9

This gene is expressed primarily in IL-1 and LPS induced neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system, including inflammatory diseases and allergies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 242 as residues: Tyr-22 to His-35.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of diseases of the immune system since expression is primarily in neutrophils, and may be useful as a growth

factor for the differentiation or proliferation of neutrophils for the treatment of neutropenia following chemotherapy.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 10

This gene is expressed primarily in activated T-cells and to a lesser extent in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune dysfunctions including cancer of the T lymphocytes and autoimmune disorders and inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of immune disorders particularly of T-cell origin and may act as a growth factor for particular subsets of T-cells such as CD4 positive cells which would make this a useful therapeutic for the treatment of HIV and other immune compromising illnesses.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 11

This gene is expressed primarily in fetal tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of many developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing fetus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor or differentiation factor for particular cell types in the developing fetus and may be useful in replacement or other types of therapy in cases where the gene is expressed aberrantly.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 12

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This gene is expressed primarily in T-cells and to a lesser extent in tumor tissue including glioblastoma, meningioma, and Wilm's tumor.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the immune system including autoimmune conditions such as rheumatoid arthritis, inflammatory disorders and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 245 as residues: Thr-9 to Ser-14.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis/ modulation of immune function disorders, including rheumatoid arthritis and inflammatory responses.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 13 30

This gene is expressed primarily in placenta and to a lesser extent in fetal liver and bone marrow.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of hematological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of

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disorders of the above tissues or cells, particularly of the hematological and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor for hematapoietic stem cells or progenitor cells in the treatment of chemotherapy patients or kidney disease.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 14

This gene is expressed primarily in stromal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of hematapoietic disorders including cancer, neutropenia, anemia, and thrombocytopenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematapoietic and immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor for hematapoietic stem cells or progenitor cells, in particular following chemotherapy treatment.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 15

The translation product of this gene shares sequence homology with epsilon-COP from Bos taurus which is thought to be important as a component of coatomer, a complex of seven proteins, that is the major component of the non-clathrin membrane coat. Preferred polypeptides encoded by this gene comprise the following amino acid sequences:

MAPPAPGPASGGSGEVDELFDVKNAFYIGSYQQCINEAXXVKLSSPERDVERD

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VFLYRAYLAQRKFGVVLDEIKPSSAPELQAVRMFADYLAHESRRDSIVAELDRE MSRSXDVTNTTFLLMAASIYLHDQNPDAALRALHQGDSLECTAMTVQILLKLD RLDLARKELKRMQDLDEDATLTQLATAWVSLATGGEKLQDAYYIFQEMADKCS PTLLLLNGQAACHMAQGRWEAAEGLLQEALDKDSGYPETLVNLIVLSQHLGKP PEVTNRYLSQLKDAHRSHPFIKEYQAKENDFDRLVLQYAPSAEAGPELSGP (SEQ ID NO:458); or RDVERDVFLYRAYLAQRKFGVVLDEIKPSSAPELQAVRMF ADYLAHESRRDSIVAELDREMSRSXDVTNTTFLLMAASIYLHDQNPDAALRALH QGDSLECTAMTVQILLKLDRLDLARKELKRMQDLDEDATLTQLATAWVSLATG GEKLQDAYYIFQEMADKCSPTLLLLNGQAACHMAQGRWEAAEGLLQEALDKD SGYPETLVNLIVLSQHLGKPPEVTNRYLSQLKDAHRSHPFIKEYQAKENDFDRL VLQYAPSA (SEQ ID NO:459).

This gene is expressed primarily in activated monocytes and T-cells, and to a lesser extent in multiple other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunomodulation, specifically relating to transport problems in these cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to epsilon-COP indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating /diagnosing problems with the cellular transport of proteins that may result in immunologic dysfunction.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 16

The translation product of this gene shares sequence homology with an RNA helicase which is thought to be important in polynucleotide metabolism. The translation product of this contig exhibits good homology to the LbeIF4A antigen of Leishmania braziliensis. The LbeIF4A antigen, or immunogenic portions of it, can be used to induce protective immunity against leishmaniasis, specifically L. donovani, L. chagasi,

L. infantum, L. major, L. braziliensis, L. panamensis, L. tropica and L. guyanensis. It can also be used diagnostically to detect Leishmania infection or to stimulate a cellular and/or humoral immune response or to stimulate the production of interleukin-12.

This gene is expressed primarily in colon cancer and to a lesser extent in pituitary.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of cancers particularly of the colon. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the gastrointestinal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 249 as residues: Glu-93 to Ala-98, Gln-150 to Leu-156, Leu-220 to Leu-231, Leu-268 to Arg-273, Val-324 to Pro-341, Arg-372 to Asn-380, Ser-405 to Gly-410, Phe-426 to Ala-433, Glu-458 to Asp-470, Arg-506 to Ser-547.

The tissue distribution and homology to RNA helicase indicates that polynucleotides and polypeptides corresponding to this gene are useful for development of diagnostic tests for colon cancer.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 17

The translation product of this contig has sequence homology to a cytoplasmic protein that binds specifically to JNK designated the JNK interacting protein-1 or JIP-1 in mice. JIP-1 caused cytoplasmic retention of JNK and inhibition of JNK-regulated gene expression.

This gene is expressed primarily in brain including pituitary cerebellum frontal cortex, fetal brain and to a lesser extent in the kidney cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of the central nervous system disorders including ischemia, epilepsy, Parkinson's disease, and schizophrenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological

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probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, 5 plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Furthermore, the translation product of this contig may suppress the effects of the JNK signaling pathway on cellular proliferation, including transformation by the Bcr-Abl oncogene. Preferred epitopes include those comprising a sequence shown in 10 SEQ ID NO: 250 as residues: Pro-6 to Ser-26, Ala-30 to Asp-41, Gly-55 to Ser-61, Gly-74 to Thr-80, Tyr-117 to Ala-123, Tyr-167 to Asp-172, Ala-212 to Cys-223, Pro-239 to Tyr-244.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for enhanced survival and/or differentiation of 15 neurons as a treatment for neurodegenerative disease.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 18

The translation product of this gene shares sequence homology with a liver stage antigen from a protozoan parasite.

This gene is expressed primarily in fetal tissue and to a lesser extent in activated T-cells and other immune cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 25 biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities and diseases of immune function. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to a protozoan antigen indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/immune modulation of parasitic infections.

# 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 19

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Preferred polypeptide encoded by this gene comprise the following polypeptide sequences:

MKAIGIEPSLATYHHIIRLFDQPGDPLKRSSFIIYDIMNELMGKRFSPKD PDDDKFFQSAMSICSSLRDLELAYQVHGLLKTGDNWKFIGPDQHRNFYYSKFF DLICLMEQIDVTLKWYEDLIPSAYFPHSQTMIHLLQALDVANRLEVIPKIWER (SEQ ID NO:460); and/or KDSKEYGHTFRSDLREEILMLMARDKHPPELQVAF ADCAADIKSAYESQPIRQTAQDWPATSLNCIAILFLRAGRTQEAWKMLGLFRKH NKIPRSELLNELMDSAKVSNSPSQAIEVVELASAFSLPICEGLTQRVMSDFAINQ EQKEALSNLTALTSDSDTDSSSDSDSDTSEGK (SEQ ID NO:461). Polynucleotides encoding such polypeptides are also provided.

This gene is expressed primarily in stromal and CD34 depleted bone marrow cells and to a lesser extent in tissues of embryonic origin.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of hematologic origin including cancers and immune dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematapoietic and immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 252 as residues: Ser-28 to Gln-34.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a growth factor for hematopoietic stem cells or progenitor cells which may be useful in the treatment of chemotherapy patients suffering from neutropenia.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 20

Preferred polypeptide fragments can be found in an alternative open reading frame. These preferred polypeptides comprise the amino acid sequence: MSSDNESDIEDEDLKLELRRLRDKHLKEIQDLQSRQKHEIESLYTKLGKVPPAVI 5 IPPAAPLSGRRRPTKSKGSKSSRSSSLGNKSPQLSGNLSGQSAASVLHPQQTL HPPGNIPESGQNQLLQPLKPSPSSDNLYSAFTSDGAISVPSLSAPGQGTSSTNTV GATVNSQAAQAQPPAMTSSRKGTFTDDLHKLVDNWARDAMNLSGRRGSKGH MNYEGPGMARKFSAPGQLCISMTSNLGGSAPISAASATSLGHFTKSMCPPQQY GFPATPFGAQWSGTGGPAPQPLGQFQPVGTASLQNFNISNLQKSISNPPGSNL 10 RTT (SEQ ID NO:462); IQDLQSRQKHEIESLYTKLGKVPPAVIIPPAAPLSGRRRR PTKSKGSKSSRSSSLGNKSPQLSGNLSGQSAASVLHPQQTLHPPGNIPESGQN QLLQPLKPSPSSDNLYSAFTSDGAISVPSLSAPGQGTSST (SEQ ID NO:463); TSDGAISVPSLSAPGQGTSSTNTVGATVNSQAAQAQPPAMTSSRKGTFTDDLH (SEQ ID NO:464); KGHMNYEGPGMARKFSAPGQLCISMTSNLGGSAPISAAS 15 ATSLGHFTK (SEQ ID NO:465); QPLKPSPSSDNLYSAFTSDGAISVPSLSAPG (SEQ ID NO:466). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in fetal liver and tissues associated with the CNS.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 20 biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver and CNS diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above 25 tissues or cells, particularly of the liver and CNS, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level 30 in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 253 as residues: Gln-26 to Lys-34.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for liver diseases such as hepatocellular carcinomas and diseases of the CNS.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 21

In an alternative reading frame, this gene shows sequence homology to two recently cloned genes, karyopherin beta 3 and Ran\_GTP binding protein 5. (See Accession Nos. gil2102696 and gnllPIDle328731.) The Ran\_GTP binding protein is related to importin-beta, the key mediator of nuclear localization signal (NLS)-dependent nuclear transport. Based on homology, it is likely that this gene may activity similar to the RAN\_GTP binding protein. Preferred polypeptide fragments comprise the amino acid sequence: VRVAAAESMXLLLECAXVRGPEYLTQMWHFMCDALIKA IGTEPDSDVLSEIMHSFAK (SEQ ID NO:467). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in thymus tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for immune disorders.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 22

This gene is expressed primarily in prostate and osteoclastoma tissues.

Preferred polypeptide fragments also comprise the amino acid sequence:

MEINNQNCFIVIDLVRTVMENGVEGLLIFGAFLPESWLIGVRCSSEPPKALLLIL

AHSQKRRLDGWSFIRHLRVHYCVSLTIHFS (SEQ ID NO:468). Also preferred are polynucleotide sequences encoding this polypeptide fragment.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, bone and prostate diseases, and cancers, particularly of the bone and prostate. Similarly, polypeptides and antibodies directed to these polypeptides are

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useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the bone and prostate systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 255 as residues: Met-1 to Ser-11.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for bone and prostate disorders, especially cancers of those systems.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 23

This gene shares sequence homology with the FK506-binding protein (FKBP-13) family, a known cytosolic receptor for the immunosuppressants. Recently, another group has cloned a very similar gene, recognizing the homology to FK506-binding protein family, calling their gene FKBP23. (See Accession No. 2827255.)

This gene is expressed primarily in lymphoid tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample, especially for those susceptible to immune suppressant therapies and for diagnosis of diseases and conditions, which include, but are not limited to, immune suppressant disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells. particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 256 as residues: Ala-19 to Val-31, Arg-38 to Gly-49, Ala-61 to Lys-66, Tyr-68 to Pro-78, Gly-116 to Ala-121, Asp-154 to Ser-162, Glu-173 to Gln-186, Phe-194 to Gly-203, Pro-207 to Val-212.

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The tissue distribution and homology to FKBP-12 and -13 indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for immune suppressant disorders.

#### 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 24

This gene is expressed primarily in the brain and in the retina. This gene maps to chromosome 8, and therefore can be used in linkage analysis as a marker for chromosome 8.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological and ocular associated disease states. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the disorders of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 257 as residues: Cys-34 to Asp-40.

The tissue distribution in retina indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and/or detection of eye disorders including blindness, color blindness, impaired vision, short and long sightedness, retinitis pigmentosa, retinitis proliferans, and retinoblastoma. Expression in the brain indicates a role in the is useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 25

This gene shows sequence homology to a newly identified class of proteins expressed in the nervous system, called stathmin family. (See Accession No. 2585991; see also Eur. J. Biochem. 248 (3), 794-806 (1997).) The stathmin family appears to be an ubiquitous phosphoprotein involved as a relay integrating various intracellular signaling pathways. These pathways affect cell proliferation and differentiation.

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Preferred polypeptide fragments comprise the amino acid sequence:

QDKHAEEVRKNKELKEEASR (SEQ ID NO:469); QQDLSPWAAPVGCPLXXASX

TCHXLPLSGCLRRQSXSLPVVAXLCFWFSCPLASLFVPGQPCVTCPFPSLPFQD

KHAEEVRKNKELKEEASR (SEQ ID NO:470). Also preferred are the
polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntintons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

# 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 26

The polynucleotide sequence of this gene contains a domain similar to a Flt3 ligand peptide. Preferred polypeptide fragments comprise the amino acid sequence: PTRCCTTQPCRSSARRPCWVPMVPSPEGREXQPTCPS (SEQ ID NO:471). Thus, this gene may have activity as binding to Flt3 receptors, a process known to promote angiogenesis and/or lymphangiogenesis.

This gene is expressed in human tonsil, and to a lesser extent in teratocarcinoma, placenta, colon carcinoma, and fetal kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the tonsil, as well as cancers, such as colon, reproductive, and kidney cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful

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in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tonsils, colon, reproductive organs, and kidneys, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 259 as residues: Pro-22 to Glu-33.

The tissue distribution in tonsil and several cancers and fetal tissues indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the tonsil or colon, such as tonsillitis, inflammatory diseases involving nose and paranasal sinuses, especially during the infection of influenza, adenoviruses, parainfluenza, rhinoviruses. The gene may also be useful in the diagnosis and treatment of neoplasms of nasopharynx or colon origins.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 27

In an alternative reading frame exists a large open reading frame that encodes a preferred polypeptide. Preferred polypeptide fragments comprise the amino acid sequence:

MKRSLNENSARSTAGCLPVPLFNQKKRNRQPLTSNPLKDDSGISTPSDNYDFP PLPTDWAWEAVNPEXAPVMKTVDTGQIPHSVSRPLRSQDSVFNSIQSNTGRSQ GGWSYRDGNKNTSLKTWXKNDFKPQCKRTNLVANDGKNSCPMSSGAQQQK QLRTPEPPNLSRNKETELLRQTHSSKISGCTMRGLDKNSALQTLKPNFQQNQY KXQMLDDIPEDNTLKETSLYQLQFKEKASSLRIISAVIESMKYWREHAQKTVLL FEVLAVLDSAVTPGPYYSKTFLMRDGKNTLPCVFYEIDRELPRLIRGRVHRCVG NYDQKKNIFQCVSVRPASVSEQKTFQAFVKIADVEMQYYINVMNET (SEQ ID NO:472); SQDSVFNSIQSNTGRSQGGWSYRDGNKNTSLKTWXKNDFKPQCKR (SEQ ID NO:473); NKETELLRQTHSSKISGCTMRGLDKNSALQTLKPNF (SEQ ID NO:474);SSLRIISAVIESMKYWREHAQKTVLLFEVLAVLDSAVTPGPYYSKTFLM (SEQ ID NO:475); and PRLIRGRVHRCVGNYDQKKNIFQCVSVRPASVSEQKT

This gene is expressed primarily in human testes.

FQAFV (SEQ ID NO:476).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, male reproductive disorders, including cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful as a hormone with reproductive or other systemic functions; contraceptive development; male infertility of testicular causes, such as Kleinfelterís syndrome, varicocele, orchitis; male sexual dysfunctions; testicular neoplasms; and inflammatory disorders such as epididymitis.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 28

This gene is expressed primarily in apoptotic T-cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases relating to T cells, as well as cancer in general. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the disorders of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for immune disorders. Moreover, since the gene was isolated from an apoptotic cell and based on the understanding of the relationship of apoptosis and cancer, it is likely that this gene may play a role in the genesis of cancer.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 29

This gene is expressed primarily in human tonsils.

Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions, which include, but are
not limited to, gastrointestinal disorders. Similarly, polypeptides and antibodies directed
to these polypeptides are useful in providing immunological probes for differential
identification of the tissue(s) or cell type(s). For a number of disorders of the above
tissues or cells, particularly of the gastrointestinal system, expression of this gene at
significantly higher or lower levels may be routinely detected in certain tissues (e.g.,
cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial
fluid or spinal fluid) or another tissue or cell sample taken from an individual having
such a disorder, relative to the standard gene expression level, i.e., the expression level
in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of gastrointestinal diseases.

# 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 30

The translation product of this gene shares sequence homology with C44C1.2 gene product of Caenorhabditis elegans with unknown function. Preferred polypeptide fragments comprise the amino acid sequence:

- GVFRPCVCGRPASLTCSPLDPEVGPYCDTPTMRTLFNLLWLALACSPVHTTLSK

  25 SDAKKAASKTLLEKSQFSDKPVQDRGLVVTDLKAESVVLEHRSYCSAKARDRH
  FAGDVLGYVTPWNSHGYDVTKVFGSKFTQISPVWLQLKRRGREMFEVTGLHD
  VDQGWMRAVRKHAKGLHIVPRLLFEDWTYDDFRNVLDSEDEIEELSKTVVQVA
  KNQHFDGFVVEVWNQLLSQKRVGLIHMLTHLAEALHQARLLALLVIPPAITPGT
  DQLGMFTHKEFEQLAPVLDGFSLMTYDYSTAHQPGPNAPLSWVRACVQVLDP
- 30 KXKWRTKSSWGSTSMXWTXRXPXDARXPVVGXRXIQXLKDHXPRMVLDSK PQ (SEQ ID NO:477); TCSPLDPEVGPYCDTPTMRTLFNLLWLALACSPVHTTLS (SEQ ID NO:478); LVVTDLKAESVVLEHRSYCSAKARDRHFAGDVLGYVTPW NSHGYDVTKVFGSKF (SEQ ID NO:479); REMFEVTGLHDVDQGWMRAVRK HAKGLHIVPRLLFEDWTYDDFRNVLDSEDE (SEQ ID NO:480); HFDGFVVEVW
- 35 NQLLSQKRVGLIHMLTHLAEALHQARLLALLVIPPATTPGTDQLGM (SEQ ID NO:481); DGFSLMTYDYSTAHQPGPNAPLSWVRACVQVLDPKXKWRTKSSW GST (SEQ ID NO:482). Also preferred are polynucleotide fragments encoding these

WO 98/54963 PCT/US98/11422

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polypeptide fragments. This gene maps to human chromosome 11, and therefore is useful in linkage analysis as a marker for chromosome 11.

This gene is expressed primarily in human T cells and to a lesser extent in human colon carcinoma.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and gastrointestinal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 263 as residues: Leu-21 to Ala-30, Ser-38 to Asp-47, Pro-87 to Asp-94, Leu-197 to Thr-204, Pro-256 to Ser-262, Thr-277 to Arg-282, Thr-293 to Trp-303.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of immune disorders and gastrointestinal diseases.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 31

The translation product of this gene shares sequence homology with Ribosomal protein L11 of Caenorhabditis elegans. (See Accession No. 156201.) Preferred polypeptide fragments comprise the amino acid sequence:

ERGVSINQFCKEFNERTKDIKEGIPLPTKILVKPDRTFEIKIGQPTVSYFLKAAAG IEKGARQTGKEVAGLVTLKHVYEIARIKAQDEAFALQDVPLSSVVRSIIGSARSL GIRVVKDLSSEELAAF QKERAIFLAAQKEADLAAQEEAAKK (SEQ ID NO:483). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in human embryo tissue and to a lesser extent in human epithelioid sarcoma and other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, development disorders and epithelial cell cancer. Similarly, polypeptides and antibodies

WO 98/54963 PCT/US98/11422

27

directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the embryonic and epithelial cell systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 264 as residues: Lys-34 to Gly-40.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of developmental disorders and epithelial cancer.

# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 32

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This gene is expressed primarily in resting T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory and general immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of disorders of immune system.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 33

This gene is believed to reside on chromosome 1. Accordingly, polynucleotides derived from this gene are useful in linkage analysis as chromosome 1 markers.

This gene is expressed primarily in prostate and to a lesser extent in soares adult brain, human umbilical vein endothelial cells, and amniotic cells.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate-related disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the urinary system and nervous system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for the diagnosis and treatment of disorders of the urinary and nervous systems.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 34

This gene shares sequence homology with R05G6.4 gene product. (See Accession No. gill326338.) This gene also shares sequence homology with the cyclophilin-like protein 20 CyP-60. (See Accession No. 1199598, see also Biochem. J. 314 (1), 313-319 (1996).) Preferred polypeptide fragments comprise the amino acid sequence: AVYTYHEKKKDTAASGYGTQNIRLSRDAVKDFDCCCLSLQPCHDPVVTPDGYL YEREAILEYILHQKKEIARQMKAYEKQRGTRREEQKELQRAASQDHVRGFLEKE SAIVSRP LNPFTAKALSGTSPDDVQPGPSVGPPSKDKDKVLPSFWIPSLTPEAK 25 ATKLEKPSRTVTCPMSGKPLRMSDLTPVHFTPLDSSVDRVGLITRSERYVCAVT RDSLSNATPCAVLRPSGAVVTLECVEKLIRKDMVDPVTGDKLTDRDIIVLORGT (SEQ ID NO:484); YLYEREAILEYILHQKKEIARQMKAYEKQRGTRREEQKELQ RAASQDHVRGFLE (SEQ ID NO:485); and FTAKALSGTSPDDVQPGPSVGPP SKDKDKVLPSFWIPSLTPEAKATKLEKPSRTVTCPMSGKPL (SEQ ID NO:486). 30 Also preferred are polynucleotide fragments that encode these polypeptide fragments.

This gene is expressed primarily in human testis and to a lesser extent in other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive disorders and in particular testicular cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing

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immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system. Expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorders of the male reproductive system and in particular of testicular cancer.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 35

The translation product of this gene shares sequence homology with Lpe5p of Saccharomyces cerevisiae which is thought to be important in the metabolism of phospholipids.

This gene is expressed primarily in liver and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, metabolic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the metabolic and nervous systems expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 268 as residues: Pro-14 to Leu-20, Lys-28 to Asn-38, Arg-109 to Arg-114, Lys-119 to Asn-124, Glu-152 to Leu-157, Pro-172 to Val-180.

The tissue distribution and homology to Lpe5p of Saccharomyces cerevisiae indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of metabolic and nervous disorders.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 36

This gene shares sequence homology with the nuclear ribonucleoprotein U (HNRNP U), encoded by *C. elegans* (See Accession gil1703576.) Preferred polypeptide fragments comprise the amino acid sequence:

5 MDTSENRPENDVPEPPMPIADQVSNDDRPEGSVEDEEKKESSLPKSFKRKISVV
SATKGVPAGNSDTEGGQPGRKRRWGASTATTQKKPSISITTESLKSLIPDIKPL
AGQEAVVDLHADDSRISEDETERNGDDGTHDKGLKICRTVTQVVPAEGQENGQ
REEEEEEKEPEAEPPVPPQVSVEVALPPPAEHEVKKVTLGDTLTRRSISQQKSGV
SITIDDPVRTAQVPSPPRGKISNIVHISNLVRPFTLGQLKELLGRTGTLVEEAFWI
10 DKIKSHCFVTYSTVEEAVATRTALHGVKWPQSNPKFLCADYAEQDELDYHRGL
LVDRPSETKTEEQGIPRPLHPPPPPPVQPPQHPRAEQREQERAVREQWAERERE
MERRERTRSEREWDRDKVREGPRSRSRSRXRRRKERAKSKEKKSEKKEKAQE
EPPAKLLDDLFRKTKAAPCIYWLPLTDSQIVQKEAERAERAKEREKRRKEQEEE
EQKEREKEAERENRQLEREKRREHSRERDRERERERDRGDRDRDRERDRE
15 RGRERDRRDTKRHSRSRSRSTPVRDRGGR (SEQ ID NO:488). Also preferred are
the polynucleotide fragments encoding this polypeptide fragments.

This gene is expressed primarily in epididymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 20 biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the male reproductive system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of 25 this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the 30 disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of male reproductive disorders.

# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 37

This gene is expressed primarily in amygdala.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory diseases and reproductive disorders. Similarly,

5 polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the amygdala, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of inflammatory diseases and reproductive disorders.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 38

This gene shares sequence homology with human opsonin protein P35 fragment. (See Accession No. R94181.) The opsonin protein activates the phagocytosis of pathogenic microbes by phagocytic cells. Preferred polypeptide fragments comprise the amino acid sequence: GCDSCPPHLPREAFAQDTQAEGECSSRAERADMCPDAP PSQEVPEGPGAAP (SEQ ID NO:489). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed in immune-related tissues such as thymus, macrophage, T cells and to a lesser extent in many other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders and infectious disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and infectious disease, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 271 as residues: Lys-9 to Arg-14, Met-38 to Asp-51.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders, as well as the treatment and/or diagnosis of infectious disease.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 39

The translation product of this gene shares sequence homology with alpha-2 type I collagen which is thought to be important in tissue repair. (See, e.g., 211607.) Preferred polypeptide fragments comprise the amino acid sequence: PQLPSCGRPW PGTASVFQSHTQGPREDPDPCRAQGSAGTHCPISLSPPRQ (SEQ ID NO:490). Also preferred are the polynucleotide sequences encoding these polypeptide sequences.

This gene is expressed primarily in the brain and to a lesser extent in the kidney and thymus

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, brain, kidney, and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, kidney, and immune disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to alpha-2 type I collagen indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of tissue repair, and brain, kidney, immune disorders.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 40

The translation product of this gene shares sequence homology with minicollagen which is thought to be important in tissue repair tumor metastasis. (See Accession No. gnllPIDld1006976.) Preferred polypeptide fragments comprise the amino acid sequence: PGFRGPSGSLGCSFFPRSLGRVLPPGCQRPGAHAD

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SSPPPTP (SEQ ID NO:491). Also preferred are polynucleotides encoding this polypeptide fragment.

This gene is expressed in ovarian cancer and to a lesser extent in dedritic cells and smooth muscle.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumor metastasis and tissue repair. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tumor metastasis and tissue repair, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 273 as residues: Asn-2 to His-11.

The tissue distribution and homology to mini-collegen gene indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of tumor metastasis and tissue repair.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 41

This gene shares sequence homology with the HIV TAT protein. (See

25 Accession No. 328416.) Preferred polypeptide fragments comprise the amino acid
sequence: EDLKKPDPASLRAASCGEGKKRKACKNCTCGLAEELEKEK
SREQMSSQPKSACGNCYLGDAFRCASCPYLGMPAFKPGEKVLLS (SEQ ID
NO:492); EDLKKPDPASLRAASCGEGKKRKACKNCTCGLAEELEKEK
SREQMSSQPKSACGNCYLGDAFRCASCPYLGMPAFKPGEKVLLSDSNLHD
30 (SEQ ID NO:493); CGNCYLGDAFRCASCPYLGMPAFKPGEKVLLSDS
(SEQ ID NO:494); SCGEGKKRKACKNCTCGLAEELEKE (SEQ ID NO:495);
SQPKSAC GNCYLGDAFRCASC (SEQ ID NO:496); and REAGQNSERQYVS
LSRD (SEQ ID NO:497). Also preferred are polynucleotide fragments encoding these
polypeptide fragments.

This gene is expressed primarily in the infant brain and to a lesser extent in the breast and testes.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, brain, testes and breast disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, testes and breast disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 274 as residues: Pro-7 to Val-15.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of brain, testes and breast, and other related disorders.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 42

This gene is expressed primarily in the infant brain, human cerebellum, and to a lesser extent in medulloblastoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, brain related disorders and medulloblastoma and other brain cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain related disorders and brain cancers, including medulloblastoma, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 275 as residues: Thr-41 to Glu-47.

WO 98/54963 PCT/US98/11422

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of human brain related disorders, brain cancers, and medulloblastoma.

## 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 43

The translation product of this gene shares sequence homology with a phosphotyrosine-independent ligand for the lck SH2 domain which is thought to be important in signal transduction related to phosphotyrosine-independent ligand for the lck SH2 domain. (See Accession No. gil1184951.) Preferred polypeptide fragments comprise the amino acid sequence: ESSGQARTLADPGPGWPRQQGMCFGSLT GLSTTPHGFLTVSAEADPRLIESLSQMLSMGFSDEGGWLTRLLQTKNYDIGAAL DTIQYSKH (SEQ ID NO:498). Also preferred are polynucleotide fragments encoding this polypeptide fragment. It is likely that this gene is a new member of a family of phosphotyrosine-independent ligands for the lck SH2 domains.

This gene is expressed primarily in the placenta and to a lesser extent in endothelial cells and neutrophil.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, reproductive, cardiovascular, immune, and infectious diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular, reproductive, and immune system, and infectious diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to a phosphotyrosine-independent ligand for the lck SH2 domain indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cardiovascular, reproductive, and immune system diseases, as well as infectious diseases.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 44

This gene is expressed primarily in the fetal brain, cerebellum and to a lesser extent in the placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal cell related disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal cell related disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 277 as residues: Thr-20 to Gly-28.

The tissue distribution and homology to proline-rich protein genes indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal cell related disorders.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 45

The translation product of this gene shares sequence homology with precerebellin of human, which is thought to be important in synaptic physiology. (See Accession No. gil180251.) It has been observed that cerebellin-like immunoreactivity is associated with Purkinje cell postsynaptic structures. Thus, it is likely that this gene also have synaptic activity. Preferred polypeptide fragments comprise the amino acid sequence: QEGSEPVLLEGECLVVCEPGRAAAGGPGGAALGEAPPGRVAFXAV RSHHHEPAGETGNGTSGAIYFDQVLVNEGGGFDRASGSFVAPVRGVYSFRFH VVKVYNRQTVQVSLMLNTWPVISAFANDPDVTREAATSSVLLPLDPGDRVSLR LRRGXSTGW (SEQ ID NO:499). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in cerebellum and infant brain. By Northern analysis, a single transcript of 2.4 kb was observed in brain tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

WO 98/54963 PCT/US98/11422

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not limited to, neuronal cell signal transduction and synaptic physiology. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal cell signal transduction and synaptic physiology expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to gene or gene family indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal cell related disorders.

## 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 46

This gene is expressed in fetal liver and spleen, and to a lesser extent in bone marrow, umbilical vein, and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the immune system, particularly hematopoiesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoiesis and immune disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 279 as residues: Asp-30 to Glu-57.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hematopieotic and immune disorders.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 47

The translation product of this gene shares sequence homology with a 12 kD nucleic acid binding protein of Feline calcivirus which is thought to be important in viral replication. (See Accession No. 59264)

This gene is expressed primarily in human cardiomyopathy and to a lesser extent in T helper cells, fetal brain and synovial sarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cardiomyopathy as well as viral infection. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 280 as residues: Trp-20 to Cys-26.

The tissue distribution in cardiomyopathy and homology to viral 12 kD nucleic acid binding protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of cardiomyopathy, including those caused by ischemic, hypertensive, congenital, valvular, or pericardial abnormalities.

The gene expression pattern may be the consequence or the cause for these conditions.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 48

The translation product of this gene shares sequence homology with tumor necrosis factor related gene product which is thought to be important in tumor necrosis, bacterial and viral infection, immune diseases and immunoreactions.

This gene is expressed primarily in colon and to a lesser extent in ovarian and breast cancers.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors of colon, ovary or breast origins. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes

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for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the colon, ovary and breast, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Tumor necrosis factors indicates that polynucleotides and polypeptides corresponding to this gene are useful for intervention of cancers of colon, ovary and breast origins, because TNF family members are known to be involved in the tumor development.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 49

The translation product of this gene shares sequence homology with mucins, such as epithelial mucin, which is thought to be important in extracellular matrix functions such as protection, lubrication and cell adhesion (See for example Accession No. R68002). Preferred polypeptide fragments comprise the following amino acid sequence: PRSRPALRPGRQRPPSHSATSGVLRPRKKPDP (SEQ ID NO:500).

Also preferred are polynucleotide fragments encoding these polypeptide fragments. Moreover, this gene maps to chromosome 22q11.2-qter, and therefore, can be used as a marker in linkage analysis for chromosome 22.

This gene is expressed primarily in corpus colosum.

Therefore, polynucleotides and polypeptides of the invention are useful as 25 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors, especially of corpus colosum, as well as metastatic lesions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell 30 type(s). For a number of disorders of the above tissues or cells, particularly of the corpus colosum and other solid tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, 35 relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to mucins indicates that polynucleotides and polypeptides corresponding to this gene are useful for serum tumor markers or immunotherapy targets because tumor cells have greatly elevated level of mucin expression and shed the molecules into the epithelial tissues.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 50

This gene is expressed primarily in CD34 depleted buffy coat cord blood and primary dendritic cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematopoietic disorders and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in CD34 depleted buffy coat cord blood and primary dendritic cells indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hematopoietic and immune disorders. Secreted or cell surface proteins in the above tissue distribution often are involved in cell activation (e.g. cytokines) or molecules involved in cell surface activation.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 51

The translation product of this gene shares sequence homology with Interferon induced 1-8 gene encoded polypeptide which is thought to be important in binding to retroviral rev responsive element. Preferred polypeptide fragment comprise the following amino acid sequences: MTLITPSXKLTFXKGNKSWSSRACSSTLVDP (SEQ ID NO:501). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

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This gene is expressed primarily in CD34 positive cells and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, retroviral infection, such as AIDS, and other immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 284 as residues: Gln-51 to Trp-62.

The tissue distribution and homology to interferon induced gene 1-8 indicates that polynucleotides and polypeptides corresponding to this gene are useful for intervention of retroviral infection including HIV. The factor may be involved in viral stability or viral entry into the cells. Alternatively, the virus/factor complex may elicit the cellular immune reaction.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 52

This gene shares sequence homology to immunoglobulin lambda chain (See Accession No. 2865484). Therefore it is likely that this gene has activity similar to an immunoglobulin lambda chain. Preferred polypeptide fragments comprise the following amino acid sequence: GHPSPALSIAPSDGSQLPCDEVPYGEAHVTRYCKKPLTNS HLETEAQSSSL (SEQ ID NO:502). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in Hodgkin's lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, Hodgkin's lymphoma and other immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 285 as residues: Pro-27 to Thr-32.

The tissue distribution in Hodgkin's lymphoma and the sequence homology indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of Hodgkin's lymphoma, since the elevated expression and secretion by the tumor mass may be indicative of tumors of this type. Additionally the gene product may be used as a target in the immunotherapy of the cancer. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 53

This gene has extensive homology to cDNA for Homo sapiens mRNA for the ISLR gene(See Accession No. AB003184). This protein is considered to be a new member of the Ig superfamily and contains a leucine-rich repeat (LRR) with conserved flanking sequences and a C2-type immunoglobulin (Ig)-like domain. These domains are important for protein-protein interaction or cell adhesion, and therefore it is possible that the novel protein ISLR may also interact with other proteins or cells. The ISLR gene was mapped on human chromosome 15q23-q24 by fluorescence in situ hybridization (See Medline Article No. 97468140). Homology to the ISLR gene has been confirmed by another independent group as well (See Accession No. Hs.102171)

This gene is expressed in a number of tissues including human retina, heart, skeletal muscle, prostate, ovary, small intestine, thyroid, adrenal cortex, testis, stomach, spinal cord, fetal lung and fetal kidney tissues, colon, tonsil and stomach cancer, and to a lesser extent in endometrial stromal cells treated with estradiol, breast tissue, synovium, lymphoma, and number of other tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors of colon, ovary and breast origins. However, due to the wide range of expression in various tissues, protein may play a vital role in the development of cancer in other tissues as well, not just those mentioned above. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the colon, ovary and breast, expression of this gene at significantly higher or lower levels may be routinely

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detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Additionally, this gene maps to chromosome 15q23-q24, and therefore, can be used as a marker in linkage analysis for chromosome 15.

The tissue distribution in tumors of colon, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 54

This gene is expressed primarily in lung, esophagus, leukemia (Jurkat cells) and breast cancers and to a lesser extent in macrophages treated with GM-CSF fetal tissues and wide range of tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer of wide range of origins. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the solid tumors, lung and leukemia, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Furthermore, due to the high expression level in lung tissue and the proposed function of the multidrug resistance protein 1 gene as the efflux pump responsible for low-drug accumulation in multidrug-resistant cells, protein as well mutants thereof, may also be beneficial as a target for gene therapy, particularly for the chronic patient. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 287 as residues: Met-1 to Lys-16.

The tissue distribution in wide range of cancers and fetal tissues indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection of cells in active proliferation, such as cancers. The gene products may be used for cancer markers or immunotherapy target.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 55

This gene maps to the X chromosome.

This gene is expressed primarily in the brain and to a lesser extent in the developing embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative disease states and developmental disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders, including sex-linked disorders, of the above tissues or cells, particularly of the neurological, developmental systems, and cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Moreover, this gene maps to the X chromosome, and therefore, may be used as a marker in linkage analysis for this chromosome.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, Klinefelter's, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental

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disorders associated with the developing embryo, sexually-linked disorders, or disorders of the cardiovascular system.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 56

5 The translation product of this gene shares sequence homology with paxillin which is thought to be important in mediating signal transduction from growth factor receptors to the cytoskeleton. Preferred polynucleotide fragments comprise the following sequence: TGGCTCACTGTCTTACAATCACTGCTGTGGAATCATGA TACCACTTTTAGCTCTTTGCATCTTCCTTCAGTGTATTTTTGTTTTTCAAGAGG 10 GGCTTGTGGTTTCAA (SEQ ID NO:506). Also preferred are polypeptide fragments encoded by these polynucleotide fragments. More preferably, polypeptide fragments comprise the amino acid sequence: LDELMAHLTEMQAKVAVRAD AGKKHLPDKQDHKASLDSMLGGLEQELQDLGIATVPKGHCASCQKPIAGKVI 15 HALGQSWHPEHFVCTHCKEEIGSSPFFERSGLXYCPNDYHQLFSPRCAYCAAP ILDKVLTAMNQTWHPEHFFCSHCGEVFGAEGFHEKDKKPYCRKDFLAMFSPK CGGCNRPVLENYLSAMDTVWHPECFVCGDCFTSFSTGSFFELDGRPFCELHYH HRRGTLCHGCGQPITGRCISAMGYKFHPEHFVCAFCLTQLSKGIFREQNDKTY CQPCFNKLF (SEQ ID NO:507); KASLDSMLGGLEQELQDLGIATVPKGHC ASCQKPIAGKVIHAL (SEQ ID NO:508); CPNDYHQLFSPRCAYCAAPILDKVL 20 TAMNQTWHPEHFFCSHCGEVFGAEG (SEQ ID NO:509); DKKPYCRKDFLAM FSPKCGGCNRPVLENYLSAMDTVWHPECFVCGDCFTSFSTGSFFELDGRPFCE L (SEQ ID NO:510); CGQPITGRCISAMGYKFHPEHFVCAFCLTQLSKGIFRE QNDKTYCQ (SEQ ID NO:511). Polynucleotide fragments encoding these preferred 25 polypeptide fragments are also contemplated.

This gene is expressed primarily in brain, and to a lesser extent in the developing embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disease states and developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or

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cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Moreover, since this gene shares homology with a gene that maps to chromosome 11, (See Accession No.T87404), gene as well as its translated product may be used for linkage analysis on chromosome 11.

The tissue distribution and homology to paxillin indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and or detection of disease states associated with abnormal signal transduction in brain and/or the developing embryo. This would include treatment or detection of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder and also in the treatment and or detection of embryonic development defects.

#### 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 57

This gene is expressed primarily in fetal spleen, brain, and to a lesser extent in six week old embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders, neurological disorders, and developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and developmental systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 290 as residues: Arg-28 to Gly-34.

The expression of this gene in fetal spleen indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/detection of immune disorders such as arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia. In addition the expression of this gene in the early embryo, indicates a key role in embryo development and hence the gene or gene product could be used in the treatment and or detection of embryonic development defects. This would include

treatment or detection of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntintons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder and also in the treatment and or detection of embryonic development defects.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 58

The translation product of this gene shares sequence homology with the gene disrupted in the neurodegenerative disease dentatorubal-pallidoluysian atrophy. Moreover a long open reading fame exists in an alternative frame. Preferred polypeptide fragments comprise the following:

- MGSSQSVEIPGGGTEGYHVLRVQENSPGHRAGLEPFFDFIVSINGSRLNKDND TLKDLLKXNVEKPVKMLIYSSKTLELRETSVTPSNLWGGQGLLGVSIRFCSFD GANENVWHVLEVESNSPAALAGLRPHSDYIIGADTVMNESEDLFSLIETHEAKP LKLYVYNTDTDNCREVIITPNSAWGGEGSLGCGIGYGYLHRIPTRPFEEGKKIS LPGQMAGTPITPLKDGFTEVQLSSVNPPSLSPPGTTGIEQSLTGLSISSTPPAVSS VLSTGVPTVPLLPPQVNQSLTSVPPMNPATTLPGLMPLPAGLPNLPNLNLNLPA PHIMPGVGLPELVNPGLPPLPSMPPRNLPGIAPLPLPSEFLPSFPLVPESSSAASS GELLSSLPPTSNAPSDPATTTAKADAASSLTVDVTPPTAKAPTTVEDRVGDSTPV SEKPVSAAVDANASESP (SEQ ID NO:512); SVEIPGGGTEGYHVLRVQENSPGH RAGLEPFFDFIVSINGSRLNKDNDTLKDLLKXNVEKPVKMLIYSSKTLELRETS
- 20 RAGLEPFFDFIVSINGSRLNKDNDTLKDLLKXNVEKPVKMLIYSSKTLELRETS
  VTPSNLWGGQGLLGVSIRFCSFDGANENVWH (SEQ ID NO:513); ESNSPAA
  LAGLRPHSDYIIGADTVMNESEDLFSLIETHEAKPLKLYVYNTDTDNCREVIITP
  NSAWGGEGSLGCGIGYGYLHRIPTRPFEEGKKISLPGQMAGTPITPLKDGFTEV
  QLSSVNPPSLSPPGTTGIEQSLTG LSISS (SEQ ID NO:514); RIPTRPFEEGKKI
- 25 SLPGQMAGTPITPLKDGFTEVQLSSVNPPSLSPPGTTGIEQSLTGLSISSTPPAVS SVLSTGVPTVPLLPPQVNQSLTSVPPMNPATTLPGLMPLPAGLPNLPNLNLNLP APHIMPGVGLPELVNPGLPPLPSMPPRN (SEQ ID NO:516); PGLPPLPSMPPRN LPGIAPLPLPSEFLPSFPLVPESSSAASSGELLSSLPPTSNAPSDPATTTAKADAA SSLTVDVTPPTAKAPTTVEDRVGDSTPVSEKPVSAAVDAN (SEQ ID NO:517).

This gene is expressed primarily in prostate cancer, and to a lesser extent in the pineal glands and in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological conditions and pulmonary disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For

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a number of disorders of the above tissues or cells, particularly of the nervous, pulmonary, and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 291 as residues: Asn-9 to Leu-14.

The abundance of this gene in the pineal gland and its homology to a gene disrupted in the neurodegenerative disease state Dentatorubral-pallidoluysian atrophy indicates that this gene may be useful in the treatment and/or detection of other neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. The abundance of this gene in fetal lung would suggest that misregulation of the expression of this protein product in the adult could lead to lymphoma or sarcoma formation, particularly in the lung; that it may also be involved in predisposition to certain pulmonary defects such as pulmonary edema and embolism, bronchitis and cystic fibrosis; and thus the gen or the gene protein encoded by the gene could be used in the detection and/or treatment of these pulmonary disorders.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 59

This gene is expressed primarily in the developing embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developmental system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

disorder.

The expression of this gene primarily in the embryo, indicates the gene plays a key role in embryo development and that the gene or the protein encoded by the gene could be used in the treatment and or detection of developmental defects in the embryo or in infants.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 60

This gene displays homology to nestin, an intermediate filament protein, the expression of which correlates with the proliferation of Central Nervous System progenitor cells and that is useful in the identification of brain tumors. This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1 (See Accession No. AA527348).

This gene is expressed primarily in kidney and to a lesser extent in brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, renal disorders and neurodegenerative conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the excretory and nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 293 as residues: Thr-128 to Asn-135.

The tissue distribution and homology to nestin indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and/or treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, its abundance in kidney indicates that it is useful in the treatment and detection of acute renal failure and other disease states associated with the kidney.

# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 61

Gene shares homology with the latrophilin-related protein 1 precursor as well as the calcium-independent alpha-latrotoxin receptor. Preferred polypeptide fragments

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comprise the following amino acid sequence:

IYKVFRHTAGLKPEVSCFENIRSCARXXXXXXXXXXXXXXXVIFGVLHVVHASVV TAYLFTVSNAFQGMFIFLFLCVLSRKIQEEYYRLFKNVPCC (SEQ ID NO:518); WIFGVLHVVHASVVTAYLFTVSNAFQGMFIFLFLCVLSRKIQEEYYRLFKNVPC

C (SEQ ID NO:519). Also preferred are polynucleotide fragments encoding these polypeptide fragments. (See Accession No. 2213659) The translation product of this gene shares sequence homology with CD 97, a seven transmembrane bound receptor.

This gene is expressed primarily in infant brain and in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders and hematopoeitic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neurological and hematopoeitic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 294 as residues: Lys-13 to Leu-21.

The tissue distribution of this gene suggest that it may be useful in the detection and/or treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder, while its expression in hematopoietic cell types indicates that the gene could be important for the treatment or detection of immune or hematopoietic disorders including arthritis, asthma and immunodeficiency diseases.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 62

This gene is expressed primarily in fetal liver and fetal spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematological and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes

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for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and hematopoetic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 295 as residues: Ser-91 to Lys-98.

The tissue distribution of this gene fetal liver and spleen indicates that the gene could be important for the treatment or detection of immune or hematopoietic disorders including arthritis, leukemia, asthma and immunodeficiency diseases.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 63

Gene shares homology with human serum amyloid protein. Preferred polypeptide fragments comprise the following amino acid sequence:
 ALTRIPPGDWVINVTAVSFAGKTTARFFHSSPPSLGDQARTDPGHQRRD (SEQ ID NO:520) (See Accession No. W13671). Also preferred are polynucleotide fragments encoding these polypeptide fragments This gene maps to chromosome 9, and therefore, may be used as a marker in linkage analysis for chromosome 9 (See Accession No. AA004342).

This gene is expressed primarily in fetal liver and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematopoietic and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution of this gene in fetal liver-spleen indicates that the gene could be important for the treatment or detection of immune or hematopoietic disorders including arthritis, leukemia, asthma, and immunodeficiency diseases.

## 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 64

This gene maps to chromosome 3, and therefore, may be used as a marker in linkage analysis for chromosome 3 (See Accession No. AA219669).

This gene is expressed specifically in the brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative disease states. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neurological systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntintons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 65

Gene shares homology with a yeast protein. Preferred polypeptide fragments comprise the following amino acid sequence: LQEVNITLPENSVWYERYKFDIP VFHL (SEQ ID NO:521). Also preferred are polynucleotide fragments encoding these polypeptide fragments. (See Accession No. 1332638)

This gene is expressed primarily in fetal tissue (fetus and fetal liver).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver disorders and cancers (e.g. hepatoblastoma). Similarly,

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polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 298 as residues: Asn-59 to Glu-64.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of liver disorders and cancers (e.g. hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells). In addition the expression in fetus would suggest a useful role for the protein product in developmental abnormalities, fetal deficiencies, pre-natal disorders and various would-healing models and/or tissue trauma.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 66

20 Gene has homology with a B-cell surface antigen which may indicate gene plays a role in the immune response, including, but not limited to disorders and infections of the immune system. Preferred polynucleotide fragments comprise the following sequence: TAGCATGTAGCCAGTCGAATAACNTATAAGGACAAAGTGGAGTC CACGCGTGCGGCCTCTAGACTAGTGGATCCCCCGGCTGCAGGATTCGGC 25 ACGAG (SEQ ID NO:523). Also preferred are polypeptide fragments encoded by these polynucleotide fragments (See Accession No.T94535). Additionally, this gene shares homology with an interferon-gamma receptor. Preferred polypeptide fragments also comprise the following amino acid sequence: MQGSGSQFRACLLCLCFSCPC SPGGPRWNSRQGGRRFPKTCRAISQNLVFKYKTFCPVRYMQPHRSSLCLHFTS 30 YVFILSTWGSLRTYSTDLKKKKKNSRGGPVPIRPKS (SEQ ID NO:522); MQGSGSQFRACLLCLCFSCPCSPGGPRWNSRQGGRRFPKTCRAISQNLVFK (SEQ ID NO:524); PVRYMQPHRSSLCLHFTSYVFILSTWGSLRTYSTDLKKKKK NSRGGPVPIRPKS (SEQ ID NO:525); and GEEQRDCSLGWRGVGMRATHCQAA RMFVLFSLPKYAGL (SEQ ID NO:526). Also preferred are polynucleotide fragments 35 encoding these polypeptide fragments

This gene is expressed primarily in T-cells and gall bladder.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological disorders and conditions (immunodeficiencies, cancer, leukemia, hematopoeisis). Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and digestive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 299 as residues: Thr-41 to Gly-52.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of immune disorders including: leukemias, lymphomas, auto-immune disorders, immuno-supressive (transplantation) and immunodeficiencies (e.g. AIDS), inflammation and hematopoeitic disorders. The expression of this gene in gall bladder would suggest a possible role for this gene product in digestive disorders, particularly of the pancreas.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 67

This gene maps to chromosome 11, and therefore, may be used as a marker in linkage analysis for chromosome 11 (See Accession No. AA011622).

This gene is expressed primarily in a variety of fetal and developmental tissues (e.g. fetal spleen, infant brain).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental, immune or neurological abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing immune and central nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 300 as residues: Ser-38 to Ser-43.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for developmental abnormalities or fetal deficiencies. The detection in infant brain would suggest a role in neurological disorders (both developmental and neurodegenerative conditions of the brain and nervous system, behavioral disorders, depression, schizophrenia, Alzheimer's disease, Parkinson's disease, Huntington's disease, mania, dementia). In addition, the detection in spleen would similarly suggest a role in detection and treatment of immunologically mediated disorders (e.g. immunodeficiency, inflammation, cancer, wound healing, tissue repair, hematopoeisis).

#### 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 68

This gene is expressed primarily in spleen, T-cells, and fetal heart.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological deficiencies, including AIDSand cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and cardiovascular systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of immune disorders including: leukemias, lymphomas, autoimmune disorders, immunodeficiencies (e.g. AIDS), immuno-suppressive conditions (transplantation) and hematopoeitic disorders. The expression in fetal heart indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of cadiovascular disorders (e.g. heart disease, restenosis, atherosclerosis, stoke, angina, thrombosis).

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 69

Gene shares homology with a human collagen protein. Preferred polypeptide fragments comprise the following amino acid sequence:

5 MPRKTSKCRQLLCSGASRNADTAARQSTCSSHRPPGKIPSLGPRRXPGCXSVP SSRGEQSTGSPAAPRCGRRDAHRGLPGGAAMTPGDTWASFNPRAGHSKSQGE GQESSGASRQDRHPVSHWVERQREAWGAPRSSSAGGVKVAATTEREPEFKIK TGKA (SEQ ID NO:527); CSGASRNADTAARQSTCSSHRPPGKIPSLGPRRXPG CXSVPSSRGEQSTGSPAAPRCGRRDAHRGLPGGAAMTPGDTWASFNPRAGHS (SEQ ID NO:528); QGEGQESSGASRQDRHPVSHWVERQREAWGAPRSSSAGG VKVAATTEREPEFKIKTGKA (SEQ ID NO:529) (See Accession No. 124886). Also preferred are polynucleotide fragments encoding these polypeptide fragments

This gene is expressed primarily in fetal heart.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 302 as residues: Pro-32 to Ser-39.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of cadiovascular disorders (e.g. heart disease, restenosis, atherosclerosis, stroke, angina, thrombosis).

## FEATURES OF PROTEIN ENCODED BY GENE NO: 70

The translation product of this gene shares sequence homology with a chicken single-strand DNA-binding protein. Preferred polypeptide fragments comprise the following amino acid sequence:

MSPRYPGGPRPILRIPNOALGGVPGSOPLLPSGMDPTROOGHPNMGGPMODI

MSPRYPGGPRPPLRIPNQALGGVPGSQPLLPSGMDPTRQQGHPNMGGPMQRM TPPRGMVPLGPQNYGGAMRPPLNALGGPGMPGMNMGPGGGRPWPNPTNAN

SIPYSSASPGNYVGPPGGGGPPGTPIMPSPADSTNSGDNMYTLMNAVPPGPNR PNFPMGPGSDGPMGGLGGMESHHMNGSLGSGDMDSISKNSPNNMSLSNOP GTPRDDGEMGGNFLNPFQSESYSPSMTMSV (SEQ ID NO:530); MSPRYPGG PRPPLRIPNQALGGVPGSQPLLPSGMDPTRQQGHPNMGGPMQRMTPPRGMVP LGPQNYGGAMRPPLNALGGPGMPGMNMGPGGGRPWPNPTNANSIPYSSASP GNY (SEQ ID. NO:531); LNALGGPGMPGMNMGPGGGRPWPNPTNANSIPYSS ASPGNYVGPPGGGGPPGTPIMPSPADSTNSGDNMYTLMNAVPPGPN (SEO ID NO:532); GPMGGLGGMESHHMNGSLGSGDMDSISKNSPNNMSLSNOPGTPR DDGEMGGNFLNPFQSESYSPSMTMSV (SEQ ID NO:533); TCEHSSEAKAFHDY (SEQ ID NO:534). Also preferred are polynucleotide fragments encoding these polypeptide fragments. (See Accession No. 1562534)

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This gene is expressed primarily in placenta and to a lesser extent in the fetal heart and a variety of other tissues and cell types.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities, fetal deficiencies, and particularly of the cardiovascular system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of developmental abnormalities or fetal deficiencies, ovarian and other endometrial cancers, reproductive dysfunction, cardiovascular disorders, and pre-natal disorders.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 71

This gene is expressed primarily in fetal liver and to a lesser extent in the breast and testes.

35 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, liver disorders (including hepatoblastomas) and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic and reproductive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and treatment of liver disorders and cancers (e.g. hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells). The expression in testes and breast indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of endocrine and reproductive disorders (e.g. sperm maturation, milk production, testicular and breast cancers).

## 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 72

This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1 (See Accession No. W93595).

This gene is expressed primarily in smooth muscle and to a lesser extent in brain.

25 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cardiovascular and neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes 30 for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cardiovascular and central nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample 35 taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of restenosis, atherosclerosis, stroke, angina, thrombosis, wound healing and other conditions of heart disease. In addition, the expression in brain would suggest that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of developmental, degenerative and behavioral conditions of the brain and nervous system (e.g. schizophrenia, depression, Alzheimer's disease, Parkinson's disease, Huntington's disease, mania, dementia, paranoia, addictive behavior and sleep disorders).

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 73

Gene shares homology with human stromalin-2. Preferred polypeptide fragments comprise the following amino acid sequence:

QAFVLLSDLLLIFSPQMIVGGRDFLRPLVFFPEATLQSELASFLMDHVFIQPGDL
GSGA (SEQ ID NO:535); ACSYLLCNPEFTFFSRADFARSQLVDLLTDRFQQE
LEELLQVG (SEQ ID NO:536),QKQLSSLRDRMVAFCELCQSCLSDVDTEIQEQV
ST (SEQ ID NO:537); QVILPALTLVYFSILWTLTHISKSDAS (SEQ ID NO:538);
STHDLTRWELYEPCCQLLQKAVDTGXVPHQV (SEQ ID NO:539). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No.R65208) This gene maps to chromosome 7, and therefore, may be used as a marker in linkage analysis for chromosome 7 (See Accession No. D52585).

This gene is expressed primarily in the brain (infant brain, adult brain, pituitary, cerebellum, hippocampus, schizophrenic hypothalmus, amygdala).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental and neurodegenerative diseases of the brain and nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those

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comprising a sequence shown in SEQ ID NO: 306 as residues: Thr-25 to Lys-36, Lys-55 to Ser-63.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and treatment of developmental, degenerative and behavioral conditions of the brain and nervous system (e.g. schizophrenia, depression, Alzheimer's disease, Parkinson's disease, Huntington's disease, mania, dementia, paranoia, addictive behavior and sleep disorders).

# FEATURES OF PROTEIN ENCODED BY GENE NO: 74

This gene is expressed primarily in the hypothalamus of a human suffering from schizophrenia.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the CNS particularly schizophrenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the CNS, such as schizophrenia expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 307 as residues: Gly-38 to Ala-44.

The tissue distribution indicates that the protein products of this gene are useful for the study, diagnosis and treatment of schizophrenia and other disorders involving the CNS.

# 30 FEATURES OF PROTEIN ENCODED BY GENE NO: 75

Preferred polypeptides of the invention comprise the following amino acid sequence encoded by this gene:

LAVSTSFICCADISTALPLGSSRPAPAPRHREHEHGHQARPPRLLXTSLMPLSTP AAAQLLWTQLTPMGGRPGGRHSPPTLHTGPRALPPGPPHPSLHVAALSLLR (SEQ ID NO:540). Polynucleotides encoding such polypeptides are also provided.

This gene is expressed primarily in endometrial tumor and to a lesser extent in amniotic cells.

WO 98/54963

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, reproductive and immune disorders particularly cancers of those systems. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 308 as residues: Ser-3 to Arg-9.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of immune and reproductive disorders particularly cancers of those systems.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 76

This gene is expressed primarily in kidney cortex and to a lesser extent in early stage human brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, renal disorders such as renal cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the kidney expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 309 as residues: Gly-38 to Gly-45, Gly-47 to Gly-52, Pro-92 to Lys-110.

The tissue distribution indicates that the protein products of this gene are useful for study, treatment and diagnosis of renal diseases such as cancer of the kidney.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 77

This gene is expressed primarily in kidney medulla.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, metabolic and renal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the metabolic and renal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for study, treatment and diagnosis of metabolic and renal diseases and disorders.

#### 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 78

This gene is expressed in chronic synovitis and microvascular endothelium.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, arthritis and atherosclerosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular and skeletal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for study, diagnosis and treatment of arthritic and other inflammatory diseases as well as cardiovascular diseases.

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PCT/US98/11422

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 79

This gene is expressed in resting T-cells and activated monocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for the study and treatment of immune diseases such as inflammatory conditions.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 80

This gene is expressed in a variety of immune system tissues, e.g., neutrophils, T-cells, and TNF induced epithelial and endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infectious and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and vascular systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 313 as residues: Met-1 to Trp-6.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of infectious diseases, immune and vascular disorders.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 81

This gene is expressed in activated neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and other immune conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of immune disorders.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 82

This gene is expressed in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory and other immune conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 315 as residues: Ala-83 to Thr-91.

The tissue distribution indicates that the protein products of this gene are useful for study and treatment of immune disorders.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 83

This gene is expressed in human neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and inflammatory system, expression of this gene, at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of disorders of the inflammatory and immune systems.

#### 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 84

This gene is expressed in human neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the inflammatory and immune systems. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the inflammatory and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of disorders of the immune and inflammatory systems.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 85

This gene is expressed in activated neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and immune system diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and inflammatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of diseases of the inflammatory and immune systems.

#### 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 86

This gene is expressed in activated neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and immune system disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the inflammatory and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 319 as residues: Met-1 to Gly-6, Gly-32 to Pro-43, Leu-55 to Gln-60.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis and treatment of disorders of the immune and inflammatory system.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 87

In specific embodiments, polypeptides of the invention comprise the sequence: EQVLALLWPRFELILEMNVQSVRSTDPQRLGGLDTRPHYITRRYAEFSSALVSIN5 QTIPNERTMQLLGQLQVEVENFVLRVAAEFSSRKEQLVFLINNYDMMLGVLME RAADDSKEVESFQQLLNARTQEFIEELLSPPFGGLVAFVKEAEALIERGQAERLR GEEARVTQLIRGFGSSWKSSVESLSQDVMRSFTNFRNGTSIIQG (SEQ ID NO:541),ALLKYRFFYQFLLGNERATAKEIRDEYVETLSKIYLSYYRSYLGRLMK VOYEEVAEKDDLMGVEDTAKKGFXSKPSRSRNTIFTLGTRGSVISPTELEAPILV 10 PHTAQR (SEQ ID NO: 542); EQRYPFEALFRSQHYXLLDNSCREYLFICEFFVVS GPXAHDLFHAVMGRTLSMTLKHLDSYLADCYDAIAVFLCIHIVLRFRNIAAKRD VPALDRYW (SEQ ID NO:543), GGLDTRPHYITRRYAEFSSALVSINQ (SEQ ID NO:544); SRKEQLVFLINNYDMMLGVL (SEQ ID NO: 545) and/or ALLKYRFFY QFLLGNERATAKEIRDEYVETLSKIYLSYYRSYLGRLMKVQYEEVAEKDDLMG VEDTAKKGFXSKPSLRSRNTIFTLGTRGSVISPTELEAPILVPHTAQRXEQRYPF 15 EALFRSQHYXLLDNSCREYLFICEFFVVSGPXAHDLFHAVMGRTLSMTLKHLD SYLADCYDAIAVFLCIHIVLRFRNIAAKRDVPALDRYWEQVLALLWPRFELILEM NVQSVRSTDPQRLGGLDTRPHYITRRYAEFSSALVSINQTIPNERTMQLLGQLQV EVENFVLRVAAEFSSRKEQLVFLINNYDMMLGVLMERAADDSKEVESFQQLLN 20 ARTQEFIEELLSPPFGGLVAFVKEAEALIERGQAERLRGEEARVTQLIRGFGSSW KSSVESLSQDVMRSFTNFRNGTS (SEQ ID NO:546). Polynucleotides encoding these polypeptides are also encompassed by the invention. The translation product of this gene shares sequence homology with suppressor of actin mutation which is thought to be important in mutation suppression.

This gene is expressed primarily in fetal liver and to a lesser extent in a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver and mutations. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the liver or cancer, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level

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in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 320 as residues: Val-53 to Arg-60, Thr-88 to Thr-94, Ala-142 to Ser-150, Gly-188 to Glu-196, Gly-208 to Ser-214, Thr-227 to Gly-232, Lys-279 to Phe-285.

The tissue distribution and homology to suppressor of actin mutation suggest that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and of liver disorder or cancer.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 88

This gene maps to chromosome 9, and therefore can be used in linkage analysis as a marker for chromosome 9. In specific embodiments, polypeptides of the invention comprise the sequence:

YEGKEFDYVFSIDVNEGGPSYKLPYNTSDDPWLTAYNFLQKNDLNPMFLDQVA KFIIDNTKGQMLGLGNPSFSDPFTGGGRYVPGSSGSSNTLPTADPFTGAGRYV PGSASMGTTMAGVDPFTGNSAYRSAASKTMNIYFPKKEAVTFDQANPTQILGK LKELNGTAPEEKKLTEDDLILLEKILSLICNSSSEKPTVQQLQILWKAINCPEDIV FPALDILRLSIKHPSVNENFCNEKEGAQFSSHLINLLNPKGKPANQLLALRTFC NCFVGQAGQKLMMSQRESLMSHAIELKSGSNKNI (SEQ ID NO: 547); HIALATLALNYSVCFHKD (SEQ ID NO: 548); HNIEGKAQCLSLISTILEVVO

- 20 DLEATFRLLVALGTLISDDSNAVQLAKS (SEQ ID NO:549); LGVDSQIKKYSS VSEPAKVSECCRFILNLL (SEQ ID NO:550); and/or YEGKEFDYVFSIDVNEGGPS YKLPYNTSDDPWLTAYNFLQKNDLNPMFLDQVAKFIIDNTKGQMLGLGNPSFS DPFTGGGRYVPGSSGSSNTLPTADPFTGAGRYVPGSASMGTTMAGVDPFTGN SAYRSAASKTMNIYFPKKEAVTFDQANPTQILGKLKELNGTAPEEKKLTEDDLI
- 25 LLEKILSLICNSSSEKPTVQQLQILWKAINCPEDIVFPALDILRLSIKHPSVNENFC NEKEGAQFSSHLINLLNPKGKPANQLLALRTFCNCFVGQAGQKLMMSQRESL MSHAIELKSGSNKNIHIALATLALNYSVCFHKDHNIEGKAQCLSLISTILEVVQD LEATFRLLVALGTLISDDSNAVQLAKSLGVDSQIKKYSSVSEPAKVSECCRFILN LL (SEQ ID NO:551). Polynucleotides encoding these polypeptides are also
- encompassed by the invention. These polypeptides share significant homology with phospholipase A2 activating protein which is thought to be important in signal transduction (see, e.g., Wang et al., Gene 161(2):237-241 (1995)).

This gene is expressed primarily in endothelial cells, to a less extent in placenta, endometrial stromal cells, osteosarcoma, testis tumor, muscle, and infant brain that are likely to be rich in blood vessles.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders in vascular system, aberrent angiogenesis, tumor angiogenesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system or tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of this gene in endothelial cells and several potential highly vascularized tissues and its homology to phospholipase A2 activating protein suggest that this gene may be involved in transducing signals for endothelial cells in angiogenesis or vasculogenesis.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 89

In specific embodiments, polypeptides of the invention comprise the sequence: YPNQDGDILRDQVLHEHIQRLSKVVTANHRALQIPEVYLREAPWPSAQSEIRTIS 20 AYKTPRDKVQCILRMCSTIMNLLSLANEDSVPGADDFVPVLVFVLIKANPPCLL STVQYISSFYASCLSGEESYWWMQFTAAVE (SEQ ID NO:552); YPNQDGDILR DQVLHEHIQRLSKVVTANHRALQIPEVYLREAPWPSAQSEIRTISAYKTPRDKVQ CILRMCSTIMNLLSLANEDSVPGADDFVPVLVFVLIKANPPCLLSTVQYISSFYA SCLSGEESYWWMQFTAAVEFIKTI (SEQ ID NO:553); YPNQDGDILRDQVL (SEQ 25 ID NO:554); EAPWPSAQSEI (SEQ ID NO:555); PVLVFVLIKANP (SEQ ID NO:560); SGEESYWWMQFTAAVEFIKTI (SEQ ID NO:556); ADDFVPVLVF VLIKANPP (SEQ ID NO:557); YKTPRDKVQCIL (SEQ ID NO:558); and/or GADDFVPVLVFVLIK (SEQ ID NO:559). The translation product of this gene shares sequence homology with human ras inhibitor and yeast VPS9p which is thought to be 30 important in golgi vacuole transport.

This gene is expressed primarily in T cells and melanocytes and to a lesser extent in a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, dysfunction and disorders involving T cells and melanocytes. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing

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immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ras inhibitor indicates that polynucleotides and polypeptides corresponding to this gene are useful for regulating signal transduction; diagnosis and treatment of disorders involving T cells and melanocytes.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 90

This gene maps to chromosome 9 and therefore polypeptides of the invention can be used in linkage analysis as a marker for chromosome 9. The translation product of this gene shares sequence homology with neuronal olfactomedin-related ER localized protein which is thought to be important in influence the maintenance, growth, or differentiation of chemosensory cilia on the apical dendrites of olfactory neurons. In specific embodiments, polypeptides of the invention comprise the sequence: SARASTQPPAGQHPGPC (SEQ ID NO:561); MPGRWRWQRDMHPARKLLSLL FLILMGTELTQD (SEQ ID NO:562); SAAPDSLLRSSKGSTRGSL (SEQ ID NO:563); AAIVIWRGKSESRIAKTPGI (SEQ ID NO:564); FRGGGTLVLPPTHT PEWLIL (SEQ ID NO:567); PLGITLPLGAPETGGGD (SEQ ID NO:565); and/or CAAETWKGSQRAGQLCALLA (SEQ ID NO:566).

This gene is expressed in pineal gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological and endocrinological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neurological or endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 323 as residues: Leu-20 to Ala-26, Arg-32 to Arg-39, Thr-104 to Gly-112.

The tissue distribution and homology to olfactomedin-related protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for maintenance, growth, or differentiation of neuron cells in pineal gland, therefore, may be useful for diagnosis and treatment of neurological disorders in pineal gland.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 91

This gene is expressed primarily in prostate and apoptotic T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate disease and T cell dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate cancer, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for detect abnormal activity in prostate and T cells or probably treatment of this abnormality.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 92

This gene is expressed primarily in prostate and to a lesser extent in smooth muscle cells, fibroblasts, and placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders in prostate or vascular system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prosate or vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain

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tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for regulating function of prostate or highly vascularized tissues, e.g. placenta.

### 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 93

This gene is expressed primarily in embryos and fetal tissues stage human and to a lesser extent in a wide variety of other proliferative tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders in embryonic development and cell proliferation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the embryonic tissues and proliferative cells, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of abnormalities in developing and proliferative cells and organs.

#### 30 FEATURES OF PROTEIN ENCODED BY GENE NO: 94

The translation product of this gene shares sequence homology with transformation related protein which is thought to be important in transformation.

This gene is expressed primarily in female reproductive tissues, i.e., breast cancer cells, placenta, and ovary and to a lesser extent in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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WO 98/54963 PCT/US98/11422

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not limited to, cancer or dysfunction of reproductive tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproduction system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 327 as residues: Ser-50 to Pro-61.

The tissue distribution and homology to transformation related protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of conditions caused by transformation, i.e. tumorigenesis in reproductive organs, e.g. breast, placenta, and ovary.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 95

This gene is expressed primarily in testes, rhabdomyosarcoma, infant brain and to a lesser extent in some tumors and highly vascularized tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumorigenesis, abnormal angiogenesis, and/or neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tumor tissues or vascular tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 328 as residues: Arg-46 to Trp-54, Pro-60 to Ile-69, Asn-116 to Ala-122, Arg-147 to Lys-153, Ser-158 to Glu-170, Ile-399 to Ser-405, Pro-486 to Met-499, Pro-502 to Asp-508.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for a range of disease states including treatment of

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tumor or vascular disorders and the treatment of neurological disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder.

### 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 96

This gene maps to chromosome 7 and therefore polynucleotides of the present invention can be used in linkage analysis as a marker for chromosome 7. The translation product of this gene is homologous to the Clostridium perfringens enterotoxin (CPE) receptor gene product and shares sequence homology with a human ORF specific to prostate and a glycoprotein specific to oligodendrocytes both of which are tissue specific proteins. (See e.g., Katahira et al., J Cell Biol. 136(6):1239-1247 (1997). PMID: 9087440; UI: 97242441.

This gene is expressed primarily in pancreas tumor and ulcerative colitis and to a lesser extent in several tumors and normal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, pancreatic disorder, ulcerative colitis, tumors and food poisoning. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system or tumorigenic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 329 as residues: Gly-147 to Met-152, Cys-177 to Lys-188.

The tissue distribution and homology to prostate and oligodendrocyte-specific protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for marker of diagnosis or treatment of disorder in pancreas, ulcerative colitis, and tumors. Furthermore, identity to the human receptor for Clostridium perfringenes entertoxin indicates that the soluble portion of this receptor could be used in the treatment of food poisoning associated with Clostridia perfringens by blocking the activity of perfringens enterotoxin.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 97

The translation product of this gene shares sequence homology with ATPase which is thought to be important in metabolism.

This gene is expressed primarily in testes and several hematopoietic cells and to a lesser extent in other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, leukemia and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 330 as residues: Leu-37 to Ala-42.

The tissue distribution and homology to ATPase indicates that polynucleotides and polypeptides corresponding to this gene are useful for marker of diagnosis and treatment of leukemia and other hematopoietic disorders.

# 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 98

In specific embodiments, polypeptides of the invention comprise the sequence:

MRSARPSLGCLPSWAFSQALNI (SEQ ID NO:568); LLGLKGLAPAEISAVCE

KGNFN (SEQ ID NO:569); VAHGLAWSYYIGYLRLILPELQARIR (SEQ ID

NO:570); TYNQHYNNLLRGAVSQRC (SEQ ID NO:571); ILLPLDCGVPDNLSM

30 ADPNIRFLDKLPQQTGDRAGIKDRVYSN (SEQ ID NO:572); SIYELLENGQRAGT

CVLEYATPLQTLFAMSQYSQAGFSGEDRLEQ (SEQ ID NO:573); AKLFCRTLE

DILADAPESQNNCRLIAYQEPADDSSFSLSQEVLRHLRQEEKEEVTVGSLKTSAV

PSTSTMSQEPELLISGMEKPLPLRTDFS (SEQ ID NO:574); and/or LLGLKGLA

PAEISAVCEKGNFNVAHGLAWSYYIGYLRLILPEL (SEQ ID NO:575).

Polynucleotides encoding these polypeptides are also encompassed by the invention.
This gene is expressed primarily in prostate BPH and to a lesser extent in bone marrow.

WO 98/54963

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, benign prostatic hypertrophy or prostate cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male urinary system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEO ID NO: 331 as residues: Ile-60 to Asn-69, Leu-106 to Asp-112, Glu-130 to Gly-136, Phe-160 to Glu-167, Pro-184 to Cys-190, Glu-197 to Ser-202, Arg-215 to Glu-221, Thr-237 to Pro-242. •

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of benign prostatic hypertrophy or prostate cancer.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 99

This gene is expressed primarily in salivary gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders or injuries of the salivary gland. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of glandular tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of disorders of, or injuries to the salivary gland or other glandular tissue.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 100

This gene maps to chromosome 15, accordingly, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 15. The translation product of this gene shares sequence homology with a *C.elegans* gene of unknown function. In specific embodiments, polypeptides of the invention comprise the sequence: DPRVRLNSLTCKHIFISLTQ (SEQ ID NO:583); TMKLLKLRRNIV KLSLYRHFTN (SEQ ID NO:576); TLILAVAASIVFIIWTTMKFRI (SEQ ID NO:577); VTCQSDWRELWVDDAIWRLLFSMILFVI (SEQ ID NO:578); MVLWR PSANNQRFAFSPLSEEEEEDEQ (SEQ ID NO:580); KEPMLKESFEGMKMRS TKQEPNGNSKVNKAQEDDL (SEQ ID NO:584); and/or KWVEENVPSSVTDVALP ALLDSDEERMITHFERSKME (SEQ ID NO:582). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in thyroid and to a lesser extent in osteoclastoma, kidney medulla, and lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, thyroid dysfunction or cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 333 as residues: Lys-107 to Leu-124, Glu-150 to Thr-159, Pro-173 to Asp-179, Ser-192 to Ser-201.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of thyroid dysfunction or cancer.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 101

This gene maps to chromosome 16, therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 16. In specific embodiments, polypeptides of the invention comprise the sequence:

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IRHELTVLRDTRPACA (SEQ ID NO:585); and/or MDFXMALIYD (SEQ ID NO:586). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in kidney cortex and to a lesser extent in adult brain, corpus colosum, hippocampus, and frontal cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment or diagnosis of neurological disorders.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 102

In specific embodiments, polypeptides of the invention comprise the sequence: MQEMMRNQDRALSNLESIPGGYNA (SEQ ID NO:587); LRRMYTDIQEPMLSA 25 AQEQF GGNPF (SEQ ID NO:588); ASLVSNTSSGEGSQPSRTENRDPLPNPWAP QT (SEQ ID NO:589); SQSSSASSGTASTVGGTTGSTASGTSGQSTTAPNLVPGV GASMFNTPG MQSLLQQITENPQLMQNMLSAPY (SEQ ID NO:590); MRSMMQSLSQNPDLAAQMMLNNPLFAGNPQLQEQMRQQLPTFLQQ (SEQ ID NO:591); MQNPDTLSAMSNPRAMQALLQIQQGLQTLATEAPGLIPGFTPGLG 30 ALGSTGGSSGTNGSNATPSENTSPTAGT (SEQ ID NO:592); TEPGHQQFI QQMLQALAGVNPQLQNPEVRFQQQLEQLSAMGFLNREANLQALIATGGDINAA IERLLGSQPS (SEQ ID NO:593); RNPAMMQEMMRNQDRALSNLESIPGGY NALRRMYTDIQEPMLSAA (SEQ ID NO:594); GNPFASLVSNTSS (SEQ ID NO:595); ENRDPLPNPWA (SEQ ID NO:595); GKILKDQDTLSQHGIHD (SEQ ID 35 NO:597); GLTVHLVIKTQNRP (SEQ ID NO:598); SELQSQMQRQLLSNPEMM (SEQ ID NO:599); PEISHMLNNPDIMR (SEQ ID NO:600); and/or RQLIMANPQMQQLIQRNP (SEQ ID NO:601). Polynucleotides encoding these

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polypeptides are also encompassed by the invention.

This gene is expressed primarily in breast.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, breast cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of tumor systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of some types of breast cancer.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 103

The translation product of this gene shares sequence homology with secreted serine proteases and lysozyme C precursor, which is thought to be important in bacteriolytic function. In specific embodiments, polypeptides of the invention comprise the sequence: NLCHVDCQDLLNPNLLAGIHCAKRIVS (SEQ ID NO:602); LDGFEGYSLSDWLCLAFVESKFN (SEQ ID NO:603);

NENADGSFDYGLFQINSHYWCN (SEQ ID NO:604); and/or NLCHVDCQDLLNPNLLAGIHCAKRIVS (SEQ ID NO:605). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infection. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 336 as residues: Ile-62 to Phe-70, Asn-78 to Asn-84.

The tissue distribution and homology to lysozyme C precursor indicates that polynucleotides and polypeptides corresponding to this gene are useful for boosting the moncyte-macrophage system and enhance the activity of immunoagents.

# 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 104

This gene is expressed primarily in apoptotic T-cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of some immune disorders.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 105

The translation product of this gene shares sequence homology with ARI

30 protein of Drosophila (accession 2058299; EMBL: locus DMARIADNE, accession X98309), which is thought to be important in axonal path-finding in the central nervous system. In specific embodiments, polypeptides of the invention comprise the sequence IREVNEVIQNPAT (SEQ ID NO:606); ITRILLSHFNWDKEKLMERYF DGNLEKLFA (SEQ ID NO:607); NTRSSAQDMPCQICYLNYPNSYF (SEQ ID NO:608); TGLECGHKFCMQCWSEYLTTKIMEEGMGQTISCPAHG (SEQ ID NO:614); CDILVDDNTVMRLITDSKVKLKYQHLITNSFVECNRLLKWCPAPD CHHVVKVQYPDAKPV (SEQ ID NO:609); CDILVDDNTVMRLITDSK

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VKLKYQHLITNSFVECNRLLKWCPAPDCHHVVKV (SEQ ID NO:610); GCNHMVCRNQNCKAEFCWVCLGPWEPHGSAWYNCNRYNEDDAKAARDAQE RSRAALQRYL (SEQ ID NO:611); FYCNRYMNHMQSLRFEHKLYAQVKQ KMEEMQQHNMSWIEVQFLKKAVDVLCQCRATLMYT (SEQ ID NO: 612);

YVFAFYLKKNNQSIIFENNQADLENATEVLSGYLERDISQDSLQDIKQKVQDKY RYCESR (SEQ ID NO:613) Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in adult brain, and to a lesser extent in endometrial tumor, melanocytes, and infant brain.

10 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases or injuries involving axonal path development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For 15 a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample 20 taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ARI protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of disease states or injuries involving axonal path development, including neurodegenerative diseases and nerve injury.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 106

The translation product of this gene shares sequence homology with cytochrome b561 [Sus scrofa] which is thought to be an integral membrane protein of neuroendocrine storage vesicles of neurotransmitters and peptide hormones.

This gene is expressed primarily in frontal cortex and to a lesser extent in rhabdomyosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to

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these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 339 as residues: Ser-18 to Pro-24.

The tissue distribution and homology to cytochrome b561 [Sus scrofa] indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of neurological disorders. This gene may also be important in regulation of some types of cancers.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 107

In specific embodiments, polypeptides of the invention comprise the sequence: MWGYLFVDAAWNFLGCLICGW (SEQ ID NO:615); MHFISSGNVSAIRSSILLL RXSLSYLGNCLRVSAIFVYFLLFLLLS (SEQ ID NO:616); and/or MDQALRGSPSE GFSTDPSPPQVGRQIPSFPPWRRLVLPKASGCFLEREWWLCVFKLRTRPGAEA HAYNSSILGGRGKGIT (SEQ ID NO:617). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in pancreas tumor and to a lesser extent in cerebellum.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, pancreatic tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred

epitopes include those comprising a sequence shown in SEQ ID NO: 340 as residues: Pro-22 to Phe-33.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of pancreatic tumors.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 108

This gene maps to chromosome 17 and therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 17. In specific embodiments, polypeptides of the invention comprise the sequence:

- MLPALASCCHFSPPEQAARLKKLQEQEKQQKVEFRKRMEKEVSDFIQDSGQIK KKFQPMNKIERSILHDVVEVAGLTSFSFGEDDDCRYVMIFKKEFAPSDEELDSY RRGEEWDPQKAEEKRNXKELAQRQ (SEQ ID NO:618); EEEAAQQGPVVV SPASDYKDKYSHLIGKGAAKDAAHMLQANKTYGCXPVANKRDTRSIEEAMNE IRAKKRLRQSGE (SEQ ID NO:619); PPRRPAQLPLTPGAGQGAGRDKAAAIRA
   HPGAPPLNHLLP (SEQ IDNO:620); AVPQAGGKQVFDLSPLELGYVRGMCVCV (SEQ ID NO:621) and/or MLPALASCCHESPPEQAARLKKLQEGEVQQKYEEDK
- (SEQ ID NO:621) and/or MLPALASCCHFSPPEQAARLKKLQEQEKQQKVEFRK
  RMEKEVSDFIQDSGQIKKKFQPMNKIERSILHDVVEVAGLTSFSFGEDDDCRYV
  MIFKKEFAPSDEELDSYRRGEEWDPQKAEEKRNXKELAQRQEEEAAQQGPVVV
  SPASDYKDKYSHLIGKGAAKDAAHMLQANKTYGCXPVANKRDTRSIEEAMNE

  10 IRAKKRLROSGE (SEQ ID NO:622) Polynycleotides encoding these polynomides
  - IRAKKRLRQSGE (SEQ ID NO:622). Polynucleotides encoding these polypeptides are also encompassed by the invention. The translation product of this gene shares sequence homology with FSA-1 which may play a role as a structural protein component of the acrosome.

This gene is expressed primarily in fetal kidney and sperm.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive disorders, especially involving acrosomal disfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

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individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 341 as residues: Glu-8 to Asn-35.

The tissue distribution and homology to FSA-1 indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of infertility due to acrosomal disfunction of sperm.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 109

This gene is expressed primarily in pituitary and to a lesser extent in epididymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 342 as residues: Met-1 to Trp-6.

Because the gene is found in both pituitary and epididymus, this indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of male reproductive disorders. This may involve a secreted peptide produced in the pituitary targeting the epididymus.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 110

In specific embodiments, polypeptides of the invention comprise the sequence: LLCPVLNSGXSWNFPHPSQPEYSFHGFHSTRLWI (SEQ ID NO:623); and/or PSTPWFLFLLGLTCPFSTSHPRWDSIPP (SEQ ID NO:624). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in resting T-cells. .

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, T-cell disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of certain immune disorders, especially those involving T-cells.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 111

This gene is expressed primarily in cerebellum and whole brain and to a lesser extent in infant brain and fetal kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 344 as residues: Asp-48 to Gly-55.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neurological disorders.

# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 112

The translation product of this gene shares sequence homology with yeast mitochondrial ribosomal protein homologous to ribosomal protein s15 of E.coli which

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is thought to be important in the early assembly of ribosomes (See Accession No. M38016). This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1.

This gene is expressed primarily in developmental tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, development of cancers and tumors in addition to healing wounds. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and developmental expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ribosomalprotein s15 of E. coli indicates that polynucleotides and polypeptides corresponding to this gene are useful for diseases related to the assembly of ribosomes in the mitochondria which is important in the translation of RNA into protein. Therefore, this indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of multiple tumors as well as in healing wounds which are thought to be under similar regulation as developmental tissues. Protein, as well as, antibodies directed against the protein have utility as tumor markers, in addition to immunotherapy targets, for the above listed tumors and tissues.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 113

The translation product of this gene shares sequence homology with human poliovirus receptor precursors which are thought to be important in viral binding and uptake. Preferred polypeptide fragments comprise the following amino acid sequence: ELSISISNVALADEGEYTCSIFTMPVRTAKSLVTVLGIPQKPIITGYKSSLREKDT ATLNCQSSGSKPAARLTWRKGDQELHGEPTRIQEDPNGKTFTVSSSVTFQVTR EDDGASIVCSVNHESLKGADRSTSQRIEVLYTPTAMIRPDPPHPREGQKLLLHC EGRGNPVPQQYLWEKEGSVPPLKMTQESALIFPFLNKSDSGTYGCTATSNMGS YKAYYTLNVND (SEQ ID NO:625). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. gnllPIDld1002627).

This gene is expressed almost exclusively in human brain tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, susceptibility to viral disease and diseases of the CNS especially cancers of that system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 346 as residues: Leu-26 to Asp-37, Lys-53 to Ser-59.

The tissue distribution and homology to poliovirus receptor precursors indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and prevention of diseases that involve the binding and uptake of virus particles for infection. It might also be helpful in genetic therapy where the goal is to insert foreign DNA into infected cells. With the help of this protein, the binding and uptake of this foreign DNA might be aided. In addition, it is expected that over expression of this gene will indicate abnormalities involving the CNS, particularly cancers of that system.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 114

The translation product of this gene shares sequence homology with YO87\_CAEEL hypothetical 28.5 KD protein ZK1236.7 in chromosome III of Caenorhabditis elegans in addition to alpha-1 collagen type III (See Accession No. gil537432). One embodiment for this gene is the polypeptide fragment(s) comprising the following amino acid sequence: VPELPDRVHQLHQAVQGCALGRPGFPGGPTH SGHHKSHPGPAGGDYNRCDRPGQVHLHNPRGTGRRGQLHPTAGPGVHRRA CPSQQLPHRLGPGVPCPSPSLTPVLPSWTQSWCG LPGYTSSS (SEQ ID NO:630). An additional embodiment is the polynucleotide fragment(s) encoding these polypeptide fragments

This gene is expressed primarily in brain cells and to a lesser extent in activated B and T cells.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegeneration and imunological disorders. Similarly, polypeptides 5 and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neural and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from 10 an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 347 as residues: Glu-34 to Glu-39, Gly-51 to Ser-72, Ala-88 to Glu-93, Gln-100 15 to Val-105.

The tissue distribution and homology to YO87\_CAEEL hypothetical 28.5 KD protein ZK1236.7 in chromosome III of Caenorhabditis elegans as well as to a conserved alpha-1 collagen type III protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons' Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorders. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 115

The translation product of this gene shares sequence homology with alpha 3 type IX collagen which is thought to be important in hyaline cartilage formation via its ability to uptake inorganic sulfate by cells (See Accession No. gil975657). One embodiment of this gene is the polypeptide fragment comprising the following amino acid sequence: SLRRPRSAAXQTLTTFLSSVSSASSSALPGSREPCDPRAPPPPR SGSAASCCSCCCSCPRRRAPLRSPRGSKRRIRQREVVDLYNGMCLQGPAGVPG RDGSPGANGIPGTPGIPGRDGFKGEKGECLRESFEESWTPNYKQCSWSSLNY GIDLGKIAECTFTKMRSNSALRVLFSGSLRLKCRNACCQRWYFTFNGAECSGP LPIEAIIYLDQGSPEMNSTINIHRTSSVEGLCEGIGAGLVDVAIWVGTCSDYPKG DASTGWNSVSRIIIEELPK (SEQ ID NO:634). An additional embodiment are the

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polynucleotide fragments encoding this polypeptide fragment.

This gene is expressed primarily in smooth muscle and to a lesser extent in synovial tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, dwarfism, spinal deformation, and specific joint abnormalities as well as chondrodysplasias i.e., spondyloepiphyseal dysplasia congenita, familial osteoarthritis. Atelosteogenesis type II, metaphyseal chondrodysplasia type Schmid and autoimmune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to alpha 3 type IX collagen indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of diseases associated with the mutation in this gene which leads to the many different types of chondrodysplasias. By the use of this product, the abnormal growth and development of bones of the limbs and spine could be routinely detected or treated in utero since the protein or muteins thereof could affect epithelial cells early in development and later the chondrocytes of the developing craniofacial structure.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 116

The translation product of this gene shares sequence homology with retrovirusrelated reverse transcriptase which is thought to be important in viral replication. One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: TKKENCRPASLMNIDTKILNKILMNQ (SEQ ID NO:640). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments (See Accession No. pirlA25313IGNHUL1).

This gene is expressed primarily in human meningima.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, retroviral diseases such as AIDS, and possibly certain cancers due to transactivation of latent cell division genes. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to retrovirus-related reverse transcriptase indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of diseases and maladies associated with retroviral infection since a functional reverse transcriptase (RT) or RT-like molecule is an integral component of the retroviral life cycle.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 117

The translation product of this gene shares sequence homology with an unknown gene from *C. elegans*, as well as weak homolog with mammalian metaxin, a gene contiguous to both thrombospondin 3 and glucocerebrosidase, is known to be required for embryonic development. Preferred polypeptide fragments comprise the following amino acid sequence: MCNLPIKVVCRANAEYMSPSGKVPXXHVGNQ VVSELGPIVQFVKAKGHSLSDGLEEVQKAEMKAYMELVNNMLLTAELYLQWC DEATVGXITHXRYGSPYPWPLXHILAYQKQWEVKRKXKAIGWGKKTLDQVLE DVDQCCQALSQRLGTQPYFFNKQPTELDALVFGHLYTILTTQLTNDELSEKVKN YSNLLAFCRRI EQHYFEDRGKGRLS (SEQ ID NO:641); MCNLPIKVVCRANAE YMSPSGKVPXXHVGNQVVSELGPIVQFVK (SEQ ID NO:642),. Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. gill326108).

This gene is expressed primarily in fetal tissues and to a lesser extent in hematopoietic cells and tissues, including spleen, monocytes, and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer; lymphoproliferative disorders; inflammation; chondrosarcoma, and Gaucher disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification

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of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and embryonic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of cancer and other proliferative disorders. Expression in embryonic tissue and other cellular sources marked by proliferating cells indicates that this protein may play a role in the regulation or cellular division. Additionally, the expression in hematopoietic cells and tissues indicates that this protein may play a role in the proliferation, differentiation, and survival of hematopoietic cell lineages. Thus, this gene may be useful in the treatment of lymphoproliferative disorders, and in the maintenance and differentiation of various hematopoietic lineages from early hematopoietic stem and committed progenitor cells.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 118

The translation product of this gene shares sequence homology with reverse transcriptase which is important in the synthesis of a cDNA chain from an RNA molecule, and is a method whereby the infecting RNA chains of retroviruses are transcribed into their DNA complements. One embodiment for this gene is the polypeptide fragment comprising the following amino acid sequence:

MXXXNSHITIFTLNVNGLNAPNERHRLANWIQSQDQVCCIQETHLTGRDTHRL

25 KIKGWRKIYQANGKQKK (SEQ ID NO:647). An additional embodiment is the polynucleotide fragments comprising polynucleotides encoding these polypeptide fragments (See Accession No. gil2072964).

This gene is expressed primarily in skin and to a lesser extent in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer, hematopoietic disorders; inflammation; disorders of immune surveillance. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the epidermis and/or hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and

WO 98/54963 PCT/US98/11422

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wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to reverse transcriptase indicates that polynucleotides and polypeptides corresponding to this gene are useful for cancer therapy. Expression in the skin also indicates that this gene is useful in wound healing and fibrosis. Expression by neutrophils also indicates that this gene product plays a role in inflammation and the control of immune surveillance (i.e. recognition of viral pathogens). Reverse transcriptase family members are also useful in the detection and treatment of AIDS.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 119

The translation product of this gene shares sequence homology with reverse transcriptase which is important in the synthesis of a cDNA copy of an RNA molecule, and is a method whereby a retrovirus reverse-transcribes its genome into an inheritable DNA copy.

This gene is expressed primarily in the frontal cortex of brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer and neurodegenerative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the CNS and peripheral nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to reverse transcriptase suggest that this is useful in the treatment of cancer and AIDS. The expression in brain indicates that it plays a role in neurodegenerative disorders and in neural degeneration.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 120

One embodiment of this gene has homology to a hypothetical protein in Schizosaccharomyces pombe (See Accession No. 2281980). Another embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: IYHLHSWIFFHFKRAFCMCFITMKVIHAHCSKLRKCXNAQISVFCTTLTASYPT (SEQ ID NO:651). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments. This gene maps to chromosome 18, and therefore, may be used as a marker in linkage analysis for chromosome 18.

This gene is expressed primarily in adult hypothalamus and to a lesser extent in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative disorders; endocrine function; and vertigo. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, CNS and peripheral nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of neurodegenerative disorders; diagnosis of tumors of a brain or neuronal origin; treatments involving hormonal control of the entire body and of homeostasis, behavioral disorders, such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental disorders associated with the developing embryo.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 121

The translation product of this gene shares sequence homology with the human IRLB protein which is thought to be important in binding to a c-myc promoter element and thus regulating its transcription (See Accession No. gil33969). This gene maps to

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chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1.

This gene is expressed primarily in brain and breast and to a lesser extent in a variety of hematopoietic tissues and cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer of the brain and breast; lymphoproliferative disorders; neurodegenerative diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the CNS, breast, and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of cancer of the brain, breast, and hematopoietic system. In addition, it may be useful for the treatment of neurodegenerative disorders, as well as disorders of the hematopoietic system, including defects in immune competency and inflammation. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 122

The translation product of this gene shares sequence homology with an ATP synthase, a key component of the proton channel that is thought to be important in the translocation of protons across the membrane.

This gene is expressed primarily in T cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, T cell lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or

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lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ATP synthase indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of defects in proton transport, homeostasis, and metabolism, as well as the diagnosis and treatment of lymphoma. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia

### FEATURES OF PROTEIN ENCODED BY GENE NO: 123

This gene maps to chromosome 15, and therefore, may be used as a marker in linkage analysis for chromosome 15.

This gene is expressed primarily in a variety of fetal tissues, including fetal liver, lung, and spleen, and to a lesser extent in a variety of blood cells, including eosinophils and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer (abnormal cell proliferation); T cell lymphomas; and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetus and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of conditions involving cell proliferation. Expression of this gene in fetal tissues, as well as in a variety of blood cell lineages indicates that it may play a role in either cellular proliferation; apoptosis; or cell survival. Thus it may be useful in the management and

treatment of a variety of cancers and malignancies. In addition, its expression in blood cells suggest that it may play additional roles in hematopoietic disorders and conditions, and could be useful in treating diseases involving autoimmunity, immune modulation, immune surveillance, and inflammation..

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 124

This gene is expressed primarily in placenta and to a lesser extent in pineal gland and rhabdomyosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental, endocrine, and female reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the [insert system where a related disease state is likely, e.g., immune], expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 357 as residues: Leu-69 to Val-76.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorders in development. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

# 30 FEATURES OF PROTEIN ENCODED BY GENE NO: 125

This gene is expressed primarily in benign prostatic hyperplasia.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of benign prostatic hyperplasia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive

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system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of benign prostatic hyperplasia. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 126

This gene is expressed primarily in apoptotic T-cells and to a lesser extent in suppressor T cells and ulcerative colitis.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases involving premature apoptosis, and immunological and gastrointestinal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of disorders involving inappropriate levels of apoptosis, especially in immune cell lineages. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases (such as AIDS), and leukemia.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 127

This gene is expressed primarily in Raji cells.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and T cell autoimmune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 360 as residues: Asp-23 to Gly-29.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of inflammation and T cell autoimmune disorders. Because the gene is expressed in cells of lymphoid origin. the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases (such as AIDS), and leukemia.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 128

25 The translation product of this gene shares sequence homology with an C. elegans coding region C47D12.2 of unknown function (See Accession No. gnllPIDle348986). One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: EDDGFNRSIHEVILKNITWY SERVLTEISLGSLLILVVIRTIQYNMTRTRDKYLHTNCLAALANMSAQFRSLHQY 30 AAQRIISLFSLLSKKHNKVLEQATQSLRGSLSSNDVPLPDYAQDLNVIEEVIRMM LEIINSCLTNSLHHNPNLVALLYKRDLFEQFRTHPSFQDIMQNIDLVISFFSSRLL QAGS (SEQ ID NO:657); EDDGFNRSIHEVILKNITWYSERVLTEISLGSLLILVV (SEQ ID NO:658); RTIQYNMTRTRDKYLHTNCLAALANMSAQFRSLHQYAAQ RIISLFSLLSKKHN (SEQ ID NO:659); KKHNKVLEQATQSLRGSLSSNDVPLPDY 35 AQD (SEQ ID NO:661); SCLTNSLHHNPNLVYALLYKRDLFEQFRTHPSFQD IMQNIDLVISFFSSRLLQAGS (SEQ ID NO:660). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments. This gene maps to

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chromosome 18, and therefore, may be used as a marker in linkage analysis for chromosome 18.

This gene is expressed primarily in smooth muscle and to a lesser extent in fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, atherosclerosis and other cardiovascular and hepatic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of circulatory system disorders such as atherosclerosis, hypertension, and thrombosis. In addition, the tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of liver disorders and cancers (e.g. hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells). In addition the expression in fetus would suggest a useful role for the protein product in developmental abnormalities, fetal deficiencies, pre-natal disorders and various would-healing models and/or tissue trauma.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 129

The translation product of this gene shares sequence homology with a ribosomal protein which is thought to be important in cellular metabolism, in addition to the *C.elegans* protein F40F11.1 which does not have a known function at the current time (See Accession No. gnllPIDle244552). Preferred polypeptide fragments comprise the following amino acid sequence:

35 MADIQTERAYQKQPTIFQNKKRVLLGETGKEKLPRVTNKNIGLGFKDT PRRLLRGTYIDKKCPFTGNVSIRGRILSGVVTQDEDAEDHCHPPRLSALHPQVQ PLREAPQEHVCTPVPL LQGRPDR (SEQ ID NO:662); MKMQRTIVIRRDYLH

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YIRKYNRFEKRHKNMSVHLSPCFRDVQIGDIVTVGECRPLSKTVRFNVLKVTK AAGTKKQFQKF (SEQ ID NO:663); MADIQTERAYQKQPTIFQNKKRVLLGET GK (SEQ ID NO:664); HCHPPRLSALHPQVQPLREAPQEHVCTPVPL LQGRPDR (SEQ ID NO:666); NIGLGFKDTPRRLLRGTYIDKKCPFTGNVSIRGRILSGVVTQ (SEQ ID NO:669); MKMQRTIVIRRDYLHYIRKYNRFEKRHKNMSVHLSP (SEQ ID NO:667); CFRDVQIGDIVTVGECRPLSKTVRFNVLKVTKAAGTKKQFQKF (SEQ ID NO:668). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in Wilm's tumor and to a lesser extent in thymus and stromal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases affecting RNA translation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Wilm's tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 362 as residues: Thr-11 to Asp-20.

The tissue distribution and homology to a ribosomal protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diseases affecting RNA translation.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 130

The translation product of this gene shares sequence homology with a yeast DNA helicase which is thought to be important in global transcriptional regulation (See Accession No. gnllPIDle243594). One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: IFYDSDWNPTVDQQA MDRAHRLGQTKQVTVYRLICKGTIEERILQRAKEKSEIQRMVISG (SEQ ID NO:670); TRMIDLLEEYMVYRKHTYXRLDGSSKISERRDMVADFQNRNDI FVFLLSTRAGGLGINLTAXDTVHF (SEQ ID NO:671); TRMIDLLEEYMVYRK HTYXRLDGSSKISERRDM (SEQ ID NO:674); RRDMVADFQNRNDIFVFLL

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STRAGGLGINLTAXDTVHF (SEQ ID NO:675), IFYDSDWNPTVDQQAMD RAHRLGQTKQVTVYRLICKG (SEQ ID NO:676); RLICKGTIEERILQRAK EKSEIQRMVISG (SEQ ID NO:678). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in amygdala.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases and disorders of the brain. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to a DNA helicase indicates that polynucleotides and polypeptides corresponding to this gene are useful for diseases affecting RNA transcription, particularly developmental disorders and healing wounds since the later are though to approximate developmental transcriptional regulation.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 131

This gene is expressed primarily in prostate and to a lesser extent in amygdala and pancreatic tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate enlargement and gastrointestinal disorders, particularly of the pancreas and gall bladder. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to

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the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of prostate diseases, including benign prostatic hyperplasia and prostate cancer. In addition, the tissue distribution in tumors of the pancreas indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tissues where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 132

This gene is expressed primarily in adult lung and to a lesser extent in hypothalamus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, pulmonary diseases and neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the pulmonary and respiratory systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of pulmonary and respiratory disorders such as emphysema, pneumonia, and pulmonary edema and emboli. In addition, the tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental

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disorders associated with the developing embryo, sexually-linked disorders, or disorders of the cardiovascular system.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 133

This gene is expressed primarily in human liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cirrhosis of the liver and other hepatic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of liver disorders such as cirrhosis, jaundice, and Hepatitus. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tissues.

#### 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 134

This gene is expressed primarily in fetal kidney and to a lesser extent in fetal liver and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, development and regeneration of liver and kidney and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive and excretory systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 367 as residues: Pro-70 to Arg-77, Tyr-102 to Thr-107.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the kidney and liver, such as cirrhosis, kidney failure, kidney stones, and liver failure, hepatoblastoma, jaundice, hepatitis, liver metabolic diseases and conditions that are attributable to the differentiation of hepatocyte progenitor cells. In addition the expression in fetus would suggest a useful role for the protein product in developmental abnormalities, fetal deficiencies, pre-natal disorders and various would-healing models and/or tissue trauma.

### 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 135

This gene is expressed primarily in brain, bone marrow, and to a lesser extent in placenta, T cell, testis and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurodegenerative and immunological diseases and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 368 as residues: Met-1 to His-6.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also

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play a role in the treatment and/or detection of developmental disorders associated with the developing embryo, or sexually-linked disorders.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 136

Translatation product of this gene is homologous to the human WD repeat protein HAN11. Preferred polypeptide fragments comprise the following amino acid sequence:

MSLHGKRKEIYKYEAPWTVYAMNWSVRPDKRFRLALGSFVEEYNNKVQLVG LDEESSEFICRNTFDHPYPTTKLMWIPDTKGVYPDLLATSGDYLRVWRVGETET RLECLLNNNKNSDFCAPLTSFDWNEVDPYLLGTSSIDTTCTIWGLETGQVLGRV NLVSGHVKTQLIAHDKEVYDIAFSRAGGGRDMFASVGADGSVRMFDLRHLEH STIIYEDPQHHPLLRLCWNKQDPNYLATMAMDGMEVVILDVRVPAHLXPGTTIE HVSMALLGPHIHPATSALQRMTTRLSSGTSSKCPEPLRTLSWPTQLXGEINNVQ WASTQPELSPSATTTAWRYSECSVGGAVPTRQGLLYFLPLPHPQS (SEQ ID

15 NO:679); MSLHGKRKEIYKYEAPWTVYAMNWSVRPDKRFRLALGSFV
EEYNNKVQLVGLDEESSEFICRNTFDHPYPTTKLMWIPDTKGVYPDLLATSGDY
LRVWRVGETETRLECLLNNNKNSDFCAPLTSFDWNEVDPYLL (SEQ ID
NO:680); SFDWNEVDPYLLGTSSIDTTCTIWGLETGQVLGRVNLVSGHVK
TQLIAHDKEVYDIAFSRAGGGRDMFASVGADGSVRMFDLRHLEHSTIIYEDPQH
40 HPLLRLCWNKODPNYLATMAMDGMEVVILDVPVPAHI VPGTTL (SEQ ID

HPLLRLCWNKQDPNYLATMAMDGMEVVILDVRVPAHLXPGTTI (SEQ ID NO:681); VGADGSVRMFDLRHLEHSTIIYEDPQHHPLLRLCWNKQDPNYLA TMAMDGMEVVILDVRVPAHLXPGTTIEHVSMALLGPHIHPATSALQRMTTRLS SGTSSKCPEPLRTLSWPTQLXGEINNVQWASTQPELSPSATTTAWRYSECSVG GAVPTRQGLLYFLPLPHPQS (SEQ ID NO:682). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in placenta, embryo, T cell and fetal lung and to a lesser extent in endothelial, tonsil and bone marrow.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological and developmental diseases in addition to cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or

cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 369 as residues: Gly-19 to Gln-28, Pro-36 to Phe-42.

The tissue distribution in tumors of colon, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

### 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 137

This gene is expressed primarily in TNF and INF induced epithelial cells, T cells and kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory conditions particularly inflammatory reactions in the kidney. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 370 as residues: Thr-67 to Gly-72, Gln-132 to Ala-145, Arg-150 to Pro-157.

The tissue distribution indicates that the protein products of this gene are useful for treating the damage caused by inflammation of the kidney.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 138

This gene maps to chromosome 1, and therefore, may be used as a marker in linkage analysis for chromosome 1 (See Accession No. D63485).

This gene is expressed primarily in breast cancer and colon cancer and to a lesser extent in thymus and fetal spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers, especially of the breast and colon tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of colon and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

# 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 139

This gene maps to chromosome 17, and therefore, can be used as a marker for linkage analysis from chromosome 17.

This gene is expressed primarily in CD34 positive cells, and to lesser extent in activated T-cells and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunologically related diseases and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and hematopoietic system, expression of this gene at significantly higher or lower levels

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may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in CD34, T-cell and neutrophils indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of hematopoietic disorders and immunologically related diseases, such as anemia, leukemia, inflammation, infection, allergy, immunodeficiency disorders, arthritis, asthma, immune deficiency diseases such as AIDS.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 140

This gene was recently cloned by another group, who called the gene KIAA0313 gene. (See Accession No. d1021609.) Preferred polypeptide fragments comprise the amino acid sequence:

- LYATATVISSPSTEXLSQDQGDRASLDAADSGRGSWTSCSSGSHDNIQTIQ HQRSWETLPFGHTHFDYSGDPAGLWASSSHMDQIMFSDHSTKYNRQNQSRES LEQAQSRASWASSTGYWGEDSEGDTGTIKRRGGKDVSIEAESSSLTSVTTEETK PVPMPAHIAVASSTTKGLIARKEGRYREPPPTPPGYIGIPITDFPEGHSHPARKP
- 20 PDYNVALQRSRMVARSSDTAGPSSVQQPHGHPTSSRPVNKPQWHKXNESDPR LAPYQSQGFSTEEDEDEQVSAV (SEQ ID NO:683); HMDQIMFSDHSTKYNRQ NQSRESLEQAQSRASWASSTGYWGE (SEQ ID NO:684); SVTTEETKPVPMP AHIAVASSTTKGLIARKEGRYREPPPTPPGYIGIPITD (SEQ ID NO:685); and VALQRSRMVARSSDTAGPSSVQQPHGHPTSSRPVNKPQW
- 25 HKXNESDPRLAPYQSQGF (SEQ ID NO:686). Also preferred are polynucleotide fragments encoding these polypeptide fragments. This gene maps to chromosome 4, and therefore, may be used as a marker in linkage analysis for chromosome 4 (See Accession No. AB002311).

This gene is expressed primarily in ovarian cancer, tumors of the Testis, brain, and colon.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, ovarian, testicle, brain and colon cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male and female reproductive systems,

WO 98/54963 PCT/US98/11422

109

expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of colon, ovary, testis, and brain origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 141

This gene is expressed primarily in spleen and colon cancer.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, colon cancer and immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the gastrointestinal trace and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of colon, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 142

Translation product is homologous to T cell translocation protein, a putative zinc finger factor (See Accession No. 340454), as well as to the G-protein coupled receptor TM5 consensus polypeptide (See Accession No. R50734). Preferred polypeptide fragments comprise the following amino acid sequence:

CLLFVFVSLGMRCLFWTIVYNVLYLKHKCNTVLLCYHLCSI (SEQ ID NO:687); ACSKLIPAFEMVMRAKDNVYHLDCFACQLCNQRXCVGDKFFLKNNXXLCQT DYEEGLMKEGYAPXVR (SEQ ID NO:688). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in fetal brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological disorders including brain cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Central Nervous System, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental disorders associated with the developing embryo.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 143

Translation product for this gene has significant homology to the Fas ligand, which is a cysteine-rich type II transmembrane protein/tumor necrosis factor receptor homolog. Mutations within this protein have been shown to result in generalized lymphoproliferative disease leading to the development of lymphadenopathy and autoimmune disease (See Medline Article No. 94185175). Preferred polypeptide

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fragments comprise the following amino acid sequence:

SALSEPGAPDRRPCPESVPRRPDDEQWPPPTALCLDVAPLPPSS (SEQ ID NO:689). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. 473565).

This gene is expressed primarily in osteoblasts, lung, and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, osteoblast-related, pulmonary, neurological, and immunological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal and nervous systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 376 as residues: Trp-33 to Thr-40, Lys-45 to Ile-63.

The tissue distribution in osteoblasts, lung, and brain combined with its homology to the Fas ligand indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. Because the Fas ligand gene is known to be expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including asthma, immune deficiency diseases such as AIDS and leukemia, and various autoimmune disorders including lupus and arthritis.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 144

This gene shares sequence homology with a 21.5 KD transmembrane protein in the SEC15-SAP4 intergenic region of yeast. (See Accession No. 1723971.) Preferred polypeptide fragments comprise the amino acid sequence:

AHASESGERWWACCGVRFGLRSIEAIGRSCCHDGPGGLVANRGRRFKWAIEL SGPGGGSRGRSDRGSGQGDSLYPVGYLDKQVPDTSVQETDRILVEKRCWDIAL

GPLKQIPMNLFIMYMAGNTISIFPTMMVCMMAWRPIQALMAISATFKMLESSSQ KFLQGLVYLIGNLMGLALAVYKCQSMGLLPTHASDWLAFIEPPERMEFSGG GLLL (SEQ ID NO:691); PVGYLDKQVPDTSVQETDRILVEKRCWDIALGPLKQ IPMNLFI (SEQ ID NO:693); and ATFKMLESSSQKFLQGLVYLIGNLMGLALAV YKCQSMGLLPTHASD (SEQ ID NO:692). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in osteoclastoma, hemangiopericytoma, liver, lung.

Therefore, polynucleotides and polypeptides of the invention are useful as 10 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, osteoclastoma, hemangiopericytoma, liver and lung tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the above tissue(s) or cell 15 type(s). For a number of disorders of the above tissues or cells, particularly of the lung and liver systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard 20 gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing osteoclastoma, hemangiopericytoma, liver and lung tumors.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 145

Translation product of this gene shares homology with the glucagon-69 gene which may indicate this gene plays a role in regulating metabolism. (See Accession No. A60318) One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

PTTKLDIMEKKKHIQIRFPSFYHKLVDSGRMRSKRETRREDSDTKHNL (SEQ ID NO:694). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in brain, kidney, colon, and testis.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, brain, kidney, colon, and testicular cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the male reproductive system, neurological, circulatory, and gastrointestinal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in tumors of brain, kidney, colon, and testis origins, indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection/treatment of neurodegenerative disease states and behavioral disorders such as Alzheimer's Disease, Parkinson's Disease, Huntingtons Disease, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder and panic disorder. In addition, the gene or gene product may also play a role in the treatment and/or detection of developmental disorders associated with the developing embryo, sexually-linked disorders, or disorders of the cardiovascular system.

# 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 146

The translation product of this gene shares sequence homology with goliath protein which is thought to be important in the regulation of gene expression during development. Protein may serve as a transcription factor. One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

- TEHIIAVMITELRGKDILSYLEKNISVQMTIAVGTRMPPKNFSRGSLVFVSISFIV LMIISSAWLIFYFIQKIRYTNARDRNQRRLGDAAKKAISKLTTRTVKKGDKETD PDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDPWLSEHCTCPMCKLNILKA LGIV (SEQ ID NO:695); TEHIIAVMITELRGKDILSYLEKNISVQMTIAVGTRMP PKNFSRGSLVFVSISFIVLM IISSAWLIFYF (SEQ ID NO:697); SISFIVLMIISSA
- 35 WLIFYFIQKIRYTNARDRNQRRLGDAAKKAISKLTTRTVKKGDKE (SEQ ID NO:698); VKKGDKETDPDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDP

WLSEHCTCPMCKLNILKALGIV (SEQ ID NO:699). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments (See Accession No. 157535). Moreover, another embodiment is the polynucleotide fragments encoding these polypeptide fragments:

- 5 MTHPGTEHIIAVMITELRGKDILSYLEKNISVQMTIAVGTRMPPKNFSRGS LVFVSISFIVLMIISSAWLIFYFIQKIRYTNARDRNQRRLGDAAKKAISKLTTRTV KKGDKETDPDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDPWLSEHCTCP MCKLNILKALGIVPNLPCTDNVAFDMERLTRTQAVNRRSALGDLAGDNSLGLE PLRTSGISPLPQDGELTPRTGEINIAVTKEWFIIASFGLLSALTLCYMIIRATASLN
- 10 ANEVEWF (SEQ ID NO:696);MTHPGTEHIIAVMITELRGKDILSYLEKNISVQM TIAVGTRMPPKNFSRGSLVFVSISFIVLMIISSAWLIFYFIQKIRYTNARDRNQRR LGDAAKKAISKLTTRT (SEQ ID NO:700); AAKKAISKLTTRTVKKGDKE TDPDFDHCAVCIESYKQNDVVRILPCKHVFHKSCVDPWLSEHCTCPMCKLNIL KALGIVPNLPC (SEQ ID NO:701); TQAVNRRSALGDLAGDNSLGLEPLRTSGI
- 15 SPLPQDGELTPRTGEINIAVTKEWFIIASFGLLSALTLCYMIIRATASLNANEVEW F (SEQ ID NO:702); PLHGVADHLGCDPQTRFFVPPNIKQWIALLQRGNCTF KEKISRAAFHNAVAVVIYNNKSKEEPVTMTHPGTEHIIAVMITELRGKDILSYLE KNISVQMTIAVGTRMPPKNFSRGSLVFVSISFIVLMIISSAWLIFYFIQKIRYTNA RDRNQRRLGDAAKKAISKLTTRTVKKGDKETDPDFDHCAVCIESYKQNDVVRI
- 20 LPCKHVFHKSCVDPWLSEHCTCPMCKLNILKALGIVPNLPCTDNVAFDMERLT RTQAVNRRSALGDLAGDNSLGLEPLRTSGISPLPQDGELTPRTGEINIAVTKEW FIIASFGLLSALTLCYMIIRATASLNANEVEWF(SEQ ID NO:703); and HGVADHLGCDPQTRFFVPPNIKQWIALLQRGNCTFKEKISRAAFHNAVAVVIY NNKSKEE (SEQ ID NO:704). An additional embodiment is the polynucleotide
- fragments encoding these polypeptide fragments. When tested against Jurkat cell lines, supernatants removed from cells containing this gene activated the GAS pathway. Thus, it is likely that this gene activates immune cells through the JAKS/STAT signal transduction pathway.

This gene is expressed primarily in macrophage, breast, kidney and to a lesser extent in synovium, hypothalamus and rhabdomyosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, schizophrenia and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and neural system, expression of this gene at

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significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to zinc finger protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of schizophrenia, kidney disease and other cancers. The tissue distribution in macrophage, breast, and kidney origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of tumors within these tissues, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 147

The translation product of this gene shares sequence homology with HNP36 protein, an equilibrative nucleoside transporter, which is thought to be important in gene transcription as well as serving as an important component of the nucleoside transport apparatus (See Accession No. 1845345). One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

- 25 MSGQGLAGFFASVAMICAIASGSELSESAFGYFITACAVIILTIICYLGLPRLEFYR
  YYQQLKLEGPGEQETKLDLISKGEEPRAGKEESGVSVSNSQPTNESHSIKAILK
  NISVLAFSVCFIFTITIGMFPAVTVEVKSSIAGSSTWERYFIPVSCFLTFNIFDWLG
  RSLTAVFMWPGKDSRWLPSWXLARLVFVPLLLLCNIKPRRYLTVVFEHDAWFI
  FFMAAFAFSNGYLASLCMCFGPKKVKPAEAETAEPSWPSSCVWVWHWGLFS
- 30 PSCSGQLCDKGWTEGLPASLPVCLLPLPSARGDPEWSGGFFF (SEQ ID NO:705); MSGQGLAGFFASVAMICAIASGSELSESAFGYFITACAVIILTIIC YLGLPRLEFYRYYQQLKLE GPGEQETKLDLISKGEEPRAGKEESGVSVSNSQ PTNESHSI (SEQ ID NO:706); SGVSVSNSQPTNESHSIKAILKNISVLAFSVCFI FTITIGMFPAVTVEVKSSIAGSSTWERYFIPVSCFLTFNIFDWLGRS (SEQ ID
- 35 NO:707),TIGMFPAVTVEVKSSIAGSSTWERYFIPVSCFLTFNIFDWLGRSLTAVF MWPGKDSRWLPSWXLARLVFVPLLLLCNIK PRRYLTVVFEHDA (SEQ ID NO:708); FGPKKVKPAEAETAEPSWPSSCVWVWHWGLFSPSCSGQLCDK

GWTEGLPASLPVCLLPLPSARGDPEWSGGFFF (SEQ ID NO:709). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in eosinophils and aortic endothelium and to a lesser extent in umbilical vein endothelial cell and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematopoietic disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to HNP36 protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of blood neoplasias and other hematopoietic disease.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 148

This gene is expressed primarily in breast cancer cell lines, thymus stromal cells, and ovary.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, endocrine and female reproductive system diseases including breast cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of endocrine disorders. In addition, the tissue distribution in tumors of thymus, ovary, and breast origins indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of these tumors, in addition to other tumors where expression has been indicated. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues

# 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 149

This gene is expressed primarily in retina and ovary and to a lesser extent in brreast cancer cell, epididymus and osteosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as 25 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal growth disorders, cancer and reproductive system disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neural and reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to 35 the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 382 as residues: Met-1 to Gly-7.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis or treatment of reproductive system disease and cancers.

# 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 150

One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

MIKDKGRARTALTSSQPAHLCPENPLLHLKAAVKEKKRNKKKKTIGSPKRIQS PLNNKLLNSPAKTLPGACGSPQKLIDGFLKHEGPPAEKPLEELSASTSGVPGLS SLQSDPAGCVRPPAPNLAGAVEFNDVKTLLREWITTISDPMEEDILQVVKYCTD LIEEKDLEKLDLVIKYMKRLMQQSVESVWNMAFDFILDNVQVVLQQTYGSTLK VT (SEQ ID NO:713); MIKDKGRARTALTSSQPAHLCPENPLLHLKAAVKE KKRNKKKKTIGSPKRIQ (SEQ ID NO:714); KRIQSPLNNKLLNSPAKT LPGACGSPQKLIDGFLKHEGPPAEKPLEELSASTSGVPGLSSLQSDPAGCVRPP

APNLAGAVEFNDVKTLLREWITTISDPM (SEQ ID NO:715);
TISDPMEEDILQVVKYCTDLIEEKDLEKLDLVIKYMKRLMQQSVE
SVWNMAFDFILDNVQVVLQQTYGSTLKVT (SEQ ID NO:716). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in 12 week embryo and to a lesser extent in hemangiopericytoma and frontal cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, growth disorders and hemangiopericytoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circular and neural system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 383 as residues: Leu-4 to Lys-11.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of growth disorders, hemangiopericytoma and other soft tissue tumors.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 151

The translation product of this gene has been found to have homology to a human DNA mismatch repair protein PMS3. Preferred polypeptide fragments comprise the following amino acid sequence: FCHDCKFPEASPAMNCEP (SEQ ID NO:717). Also preferred are polynucleotide fragments encoding these polypeptide fragments (See Accession No. R95250).

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, lymphoma, immunodeficiency diseases, and cancers resulting from genetic instability. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 384 as residues: Met-1 to Lys-6.

The tissue distribution in neutrophils and the sequence homology indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of Hodgkin's lymphoma, since the elevated expression and secretion by the tumor mass may be indicative of tumors of this type. Additionally the gene product may be used as a target in the immunotherapy of the cancer. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia. Furthermore, its homology to a known DNA repair protein would suggest gene may be useful in establishing cancer predisposition and prevention in gene therapy applications.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 152

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infectious diseases and lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of inflammation and infectious diseases.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 153 15

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One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence:

MASSVPAGGHTRAGGIFLIGKLDLEASLFKSFQWLPFVLRKKC NFFCWDSSAHSLPLHPLSASCSAPACHASDTHLLYPSTRALCPSIFAWLVAPHS VFRTNAPGPTPSSQSSPVFPVFPVSFMALIVCXLVCC (SEQ ID NO:720); MASSVPAGGHTRAGGIFLIGKLDLEASLFKSFQWLPFVLRKKCNFFCWDSSAH SLPLHPLSASCSAPACHA (SEQ ID NO:721);FAWLVAPHSVFRTNAPGPTPS SQSSPVFPVFPVSFMALIVCXLVCC (SEQ ID NO:722). An additional embodiment is the polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammation and infectious disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred

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epitopes include those comprising a sequence shown in SEQ ID NO: 386 as residues: Ser-11 to Pro-17.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment of infectious diseases and inflammation.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 154

This gene is expressed in multiple tissues including ovary, uterus, adipose tissue, brain, and the liver.

10 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, uterine, ovarian, brain, and liver cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes 15 for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the female reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution of this gene indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnostic or therapeutic uses in the treatment of the female reproductive system, obesity, and liver disorders, particularly cancer in the above tissues.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 155

This gene maps to chromosome 3, and therefore, may be used as a marker in linkage analysis for chromosome 3 (See Accession No. D87452).

This gene is expressed in multiple tissues including brain, aortic endothelial cells, smooth muscle, pituitary, testis, melancytes, spleen, nertrophils, and placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological disorders including immunodeficiencies, cancers of the brain and the female reproductive system, as well as cardiovascular disorders, such as atherosclerosis and stroke. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution suggest that polynucleotides and polypeptides corresponding to this gene are useful in treatment/detection of disorders in the nervous system, including schizophrenia, neurodegeneration, neoplasia, brain cancer as well as cardiovascular and female reproductive disorders including cancer within the above tissues.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 156

The translation product of this gene shares sequence homology with the human gene encoding cytochrome b561 (See Accession No. P10897). Cytochrome b561 is a transmembrane electron transport protein that is specific to a subset of secretory vesicles containing catecholamines and amidated peptides. This protein is thought to supply reducing equivalents to the intravesicular enzymes dopamine-beta-hydroxylase and alpha-peptide amidase. Preferred polypeptides of the invention comprise the amino acid sequence:

MAMEGYWRFLALLGSALLVGFLSVIFALVWVLHYREGLGWDGSALEFNWHP VLMVTGFVFIQGIAIIVYRLPWTWKCSKLLMKSIHAGLNAVAAILAIISVVAVFE NHNVNNIANMYSLHSWVGLIAVICYLLQLLSGFSVFLLPWAPLSLRAFLMPIHV YSGIVIFGTVIATALMGLTEKLIFSLRDPAYSTFPPEGVFVNTLGLLILVFGALIF WIVTRPQWKRPKEPNSTILHPNGGTEQGARGSMPAYSGNNMDKSDSEL NSEVAARKRNLALDEAGQRSTM (SEQ ID NO:724); as well as antigenic fragments of at least 20 amino acids of this gene and/or biologically active fragments. Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in anergic T-cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune system and metabolism related diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological

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probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product or RNA of this gene is useful for treatment or diagnosis of immune system and metabolic diseases or conditions including Tay-Sachs disease, phenylketonuria, galactosemia, various porphyrias, and Hurler's syndrome.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 157

The translation product of this gene shares sequence homology with collagen which is important in mammalian development. This gene also shows sequence homology with bcl-2. (See Accession No. P80988.) Preferred polypeptide fragments comprise the amino acid sequence: PGRAGPSPGLSLQLPAEPGHPAGNLAPL TSRPQPLCRIPAVPG (SEQ ID NO:725). Also preferred are polynucleotide sequences encoding this polypeptide fragment.

This gene is expressed primarily in HL-60 tissue culture cells and to a lesser extent in liver, breast, and uterus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunological diseases, hereditary disorders involving the MHC class of immune molecules, as well as developmental disorders and reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and reproductive system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those

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comprising a sequence shown in SEQ ID NO: 390 as residues: Ser-39 to Gly-46, Leu-49 to Ala-62.

The tissue distribution and homology to collagen indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hereditary MHC disorders and particularly autoimmune disorders including rheumatoid arthritis, lupus, scleroderma, and dermatomyositis, as well as many reproductive disorders, including cancer of the uterus, and breast tissues.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 158

This gene is expressed primarily in the amygdala region of the brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, a variety of brain disorders, particularly those effecting mood and personality. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and/or diagnosis of a variety of brain disorders, particularly bipolar disorder, unipolar depression, and dementia.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 159

This gene is expressed in a variety of tissues and cell types including brain, smooth muscle, kidney, salivary gland and T-cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers of a variety of organs including brain, smooth muscle, kidney, salivary gland and T-cells and cardiovascular disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

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of the above tissues or cells, particularly of the central nervous, urinary, saliyary, digestive, and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in brain, smooth muscle, and T-cells indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of various neurological, and cardiovascular disorders, but not limited to cancer within the above tissues. Additionally the gene product may be used as a target in the immunotherapy of the cancer. Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 160

The translation product of this gene shares sequence homology with collagen which is thought to be important in cellular interactions, extracellular matrix formation. and has been found to be an identifying determinant in autoimmune disorders. Moreover, this gene shows sequence homology with the yeast protein, Sls1p, an endoplasmic reticulum component, involved in the protein translocation process in Yeast Yarrowia lipolytica. (See Accession No. 1052828; see also J. Biol. Chem. 271, 11668-11675 (1996).) With mouse, this same region shows sequence homology with the heavy chain of kinesin. (See Accession No. 2062607.) Recently, suppression of the heavy chain of kinesin was shown to inhibits insulin secretion from primary cultures of mouse beta-cells. (See Endocrinology 138 (5), 1979-1987 (1997).) Moreover, kinesin was found associated with drug resistance and cell immortalization. (See 468355.) Thus, it is likely that this gene also act as a genetic suppressor elements.

This gene is expressed primarily in the greater omentum and to a lesser extent in a variety of organs and cell types including gall bladder, stromal bone marrow cells, lymph node, liver, testes, pituitary, and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders of the endocrine, gastrointestinal, and immunological systems, including autoimmune disorders and cancers in a variety of organs and cell types.

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Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and gastrointestinal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 393 as residues: Asn-27 to Leu-47, Gln-81 to Lys-88, Asp-93 to Lys-102, Asn-107 to Leu-116, Met-129 to Glu-141, Glu-150 to Asp-157, Lys-176 to Glu-185, Glu-333 to Tyr-349, Cys-393 to Leu-403, Gln-423 to Gly-429.

The tissue distribution in within various endocrine and immunological tissues combined with the sequence homology to a conserved collagen motif indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis of various autoimmune disorders including, but not limited to, rheumatoid arthritis, lupus erthyematosus, scleroderma, dermatomyositis Because the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 161

This gene has homology to the tissue inhibitor of metalloproteinase 2. Such inhibitors are vital to proper regulation of metalloproteins such as collagenases (See Accession No. P16368). In addition, this gene maps to chromosome 17, and therefore, may be used as a marker in linkage analysis for chromosome 17 (See Accession No. P16368).

This gene is expressed primarily in several types of cancer including osteoclastoma, chondrosarcoma, and rhabdomyosarcoma and to a lesser extent in several non-malignant tissues including synovium, amygdala, testes, placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, various types of cancer, particularly cancers of bone and cartilage, as well as various autoimmune disorders. Similarly, polypeptides and antibodies directed

to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the musculoskeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in various cancers and the sequence homology to a collagenase inhibitor indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection of various autoimmune disorders such as rheumatoid arthritis, lupus, scleroderma, and dermatomyositis. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 162

This gene is homologous to the mitochondrial ATP6 gene and therefore is likely a homolog of this gene family (See Accession No. X76197).

This gene is expressed primarily in brain tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, a variety of brain disorders, including Down's syndrome, depression, Schizophrenia, and epilepsy. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in brain tissue indicates this gene is useful for diagnosis of various neurological disorders including, but not limited to, brain cancer. Additionally the gene product may be used as a target in the immunotherapy of cancer in the brain as well as for the diagnosis of metabolic disorders such as obesity Tay-Sachs disease, phenylketonuria and Hurler's Syndrome.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 163

This gene is expressed primarily in placenta, neutrophils, and microvascular endothelial cells and to a lesser extent in multiple tissues including brain, prostate, spleen, thymus, and bone.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neutropenea and other diseases of the immune system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in placenta indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis various female reproductive disorders. Additionally the gene product may be used as a target in the immunotherapy of various cancers. Because the gene is expressed in some cells of lymphoid and endocrine origin, the natural gene product may be involved in immune functions and metabolism regulation, respectively. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 164

This gene is expressed primarily in neutrophils, monocytes, bone marrow, and fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune system disorders including, but not limited to, autoimmune disorders such as lupus, and immunodeficiency disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

WO 98/54963

129

of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in various immune system tissue indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis of various immunological disorders such as Hodgkin's lymphoma, arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 165

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The translation product of this gene shares sequence homology with dystrophin which is thought to be defective in both Duchene and Becker Muscular Dystrophy. 15 Preferred polypeptide fragments comprise the following amino acid sequence: MKLLGECSSSIDSVKRLEHKLKEEEESLPGFVNLHSTETQTAGVIDRWELLQAQ ALSKELRMKQNLQKWQQFNSDLNSIWAWLGDTEEELEQLQRLELSTDIQTIELO IKKLKELQKAVDHRKAIILSINLCSPEFTQADSKESRDLQDRLXQMNGRWDRV CSLLEEWRGLLQDALMQCQGFHEMSHGLLLMLENIDRRKNEIVPIDSNLDAEIL 20 QDHHKQLMQIKHELLESQLRVASLQDMSCQLLVNAEGTDCLEAKEKVHVIGNR LKLLLKEVSRHIKELEKLLDVSSSQQDLSSWSSADELDTSGSVSPXSGRSTPNR QKTPRGKCSLSQPGPSVSSPHSRSTKGGSDSSLSEPXPGRSGRGFLFRVLRAA LPLQLLLLLIGLACLVPMSEEDYSCALSNNFARSFHPMLRYTNGPPPL (SEQ ID NO:726); MKLLGECSSSIDSVKRLEHKLKEEEESLPGFVNLHSTETQTAGVIDR 25 WELLQAQALSKELRMKQNLQKWQQFNSDLNSIWAWLGDTEEELEQLQRLELS TDIQTIELQIK (SEQ ID NO:727); KLKELQKAVDHRKAIILSINLCSPEFTQADSK ESRDLQDRLXQMNGRWDRVCSLLEEWRGLLQDALMQCQGFHEMSHGLLLML ENIDRRKNEIVPIDSNLDAEILQDHHKQLMQIKHELLESQLRVASLQDMSCQL (SEQ ID NO:728); QDMSCQLLVNAEGTDCLEAKEKVHVIGNRLKLLLKEVS 30 RHIKELEKLLDVSSSQQDLSSWSSADELDTSGSVSPXSGRSTPNRQKTPRGKCS LSQPGPSVSSPHS (SEQ ID NO:729); DSSLSEPXPGRSGRGFLFRVLRAAL PLQLLLLLIGLACLVPMSEEDYSCALSNNFARSFHPMLRYTNGPPPL (SEQ ID NO:730). Also preferred are polynucleotide fragments encoding these polypeptide fragments. Furthermore, this gene maps to chromosome 6, and therefore, may be used

This gene is expressed in numerous tissues including the heart, kidney, and brain.

as a marker in linkage analysis for chromosome 6 (See Accession No. N62896).

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, musculoskeletal disorders including Muscular Dystrophy and cardiovascular diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the muscle tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to dystrophin indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of Muscular Dystrophy and other muscle disorders.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 166

This gene is expressed primarily in human cerebellum.

20 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the central nervous system, including Alzheimer's Disease, Parkinson's Disease, ALS, and mental illnesses. Similarly, polypeptides and antibodies 25 directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, 30 synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEO ID NO: 399 as residues: Pro-20 to Gly-26, Leu-37 to Pro-42, His-57 to Gly-63.

The tissue distribution indicates that the protein products of this gene are useful for treatment/diagnosis of diseases of the central nervous system and may protect or

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enhance survival of neuronal cells by slowing progression of neurodegenerative diseases.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 167

5 Preferred polypeptides encoded by this gene comprise the following amino acid sequence:

MKLLICGNYLAPSHSESSRRCCLLCFYPLCLEINFGMKVFLSMPFLVLFQ SLIQED (SEQ ID NO:731). Polynucleotides encoding such polypeptides are also provided. This gene is believed to reside on chromosome 15. Therefore polynucleotides derived from this gene are useful in linkage analysis as chromosome 15 markers.

This gene is expressed primarily in human testes tumor and to a lesser extent in normal human testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are 15 not limited to, diseases of the testes, particularly cancer, and other reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of 20 the male reproductive tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily 25 fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for treatment/diagnosis of testicular diseases including cancers.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 168

This gene is expressed primarily in fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, conditions affecting hematopoietic development and metabolic diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the

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hepatic system, and fetal hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 401 as residues: His-7 to Trp-17, Leu-19 to Lys-27, Pro-33 to Gly-44, Lys-68 to Gly-74, Lys-85 to Cys-95.

The tissue distribution indicates that the protein products of this gene are useful for treatment/diagnosis of diseases of the developing liver and hematopoietic system, and act as a growth differentiation factor for hematopoietic stem cells.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 169

The polypeptide encoded by this gene is believed to be a membrane bound receptor. The extracellular domain of which is expected to consist of the following amino acid sequence:

RILLVKYSANEENKYDYLPTTVNVCSELVKLVFCVLVSFCVIKKDHQSRNLKY ASWKEFSDFMKWSIPAFLYFLDNLIVFYVLSYLQPAMAVIFSNFSIITTALLFRIV LKXRLNWIQWASLLTLFLSIVALTAGTKTLQHNLAGRGFHHDAFFSPSNSCLL

- FRNECPRKDNCTAKEWTFPEAKWNTTARVFSHIRLGMGHVLIIVQCFISSMANI YNEKILKEGNQLTEXIFIQNSKLYFFGILFNGLTLGLQRSNRDQIKNCGFFYGH S (SEQ ID NO:732). Thus, preferred polypeptides encoded by this gene comprise the extracellular domain as shown above. It will be recognized, however, that deletions of either end of the extracellular domain up to the first cysteine from the N-terminus and the first cysteine of the C-terminus, is expected to retain the biological functions of the full-length extracellular domain because the cysteines are thought to be responsible for providing secondary structure to the molecule. Thus, deletions of one or more amino
- acids from either end (or both ends) of the extracellular domain are contemplated. Of course, further deletions including the cysteines are also contemplated as useful as such polypeptides is expected to have immunological properties such as the ability to evoke and immune response. Polynucleotides encoding all of the foregoing polypeptides are provided.

This gene is expressed primarily in human osteoclastoma and to a lesser extent in hippocampus and chondrosarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

not limited to, cancers, particularly those of the bone and connective tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 402 as residues: Met-1 to Cys-6, Ala-41 to Tyr-49, Lys-76 to Lys-84.

The tissue distribution indicates that the protein products of this gene are useful for diagnosis of cancers of the bone and connective tissues, and may act as growth factors for cells involved in bone or connective tissue growth.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 170

Preferred polypeptides encoded by this gene comprising the following amino acid sequence:

NSVPNLQTLAVLTEAIGPEPAIPRXPREPPVATSTPATPSAGPQPLPTGTV LVPGGPAPPCLGEAWALLLPPCRPSLTSCFWSPRPSPWKETGV (SEQ ID NO:733). Polynucleotides encoding such polypeptides are also provided herein.

This gene is expressed primarily in hematopoietic progenitor cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the blood including cancer and autoimmune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the blood/circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 403 as residues: Gln-4 to His-10, Pro-25 to His-32.

WO 98/54963 PCT/US98/11422

134

The tissue distribution indicates that the protein products of this gene are useful for diagnosis of diseases involving growth differentiation of hematopoietic cells.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 171

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Preferred polypeptides encoded by this gene comprise the following amino acid sequences: ALQLAFYPDAVEEWLEENVHPSLQRLQXLLQDLSEVSAPP (SEQ ID NO:734); and/or CHPPALAGTLLRTPEGRAHARGLLLEAGGA (SEQ ID NO:735). Polynucleotides encoding such polypeptides are also provided. The protein product of this gene shares sequence homology with metallothionines. Thus, polypeptide encoded by this gene are expected to have metallothionine activity, such activities are known in the art and described elsewhere herein.

This gene is expressed primarily in kidney cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases of the kidney including cancer and renal dysfunction. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 404 as residues: Ser-47 to Gln-52.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of diseases of the kidney including kidney failure.

FEATURES OF PROTEIN ENCODED BY GENE NO: 172

This gene is expressed primarily in 12 week old early stage human.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for

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differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing embryo, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 405 as residues: Gln-31 to Thr-43, Gly-51 to Ser-58, Pro-65 to Pro-72.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of developmental problems with fetal tissue. The gene may be involved in vital organ development in the early stage, especially hematopoiesis, cardiovascular system, and neural development.

### 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 173

The translation product of this gene shares sequence homology with TGN38, an integral membrane protein previously shown to be predominantly localized to the trans-Golgi network (TGN) of cells.

This gene is expressed primarily in developing embryo and to a lesser extent in cancer tissues including lymphoma, endometrial, protate and colon.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the developing fetus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 406 as residues: His-65 to Ser-72, Pro-82 to Gly-91, Pro-98 to Glu-118, Ser-126 to Gly-166, Pro-180 to Asp-188, Tyr-209 to Lys-214, Gln-220 to Leu-228.

The tissue distribution and homology to an integral membrane protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for

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diagnosis of cancers and developmental abnormalities where aberrant expression relates to an abnormality.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 174

The translation product of this gene shares sequence homology with a dnaJ heat shock protein from E. coli which is allelic to sec63, a gene that affects transit of nascent secretory proteins across the endoplasmic reticulum in yeast.

This gene is expressed primarily in Hodgkin's lymphoma and to a lesser extent in testes.

10 Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells. 15 particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to 20 the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 407 as residues: Thr-13 to Trp-21, Arg-74 to Asp-81.

The tissue distribution and homology to dnaJ indicates that polynucleotides and polypeptides corresponding to this gene are useful as a diagnostic for cancer including Hodgkin's lymphoma.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 175

This gene is expressed primarily in endothelial cells and to a lesser extent in bone marrow stromal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, diseases involving angiogenic abnormalities including diabetic retinopathy, macular degeneration, and other diseases including arteriosclerosis and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell

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type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for treating diseases where an increase or decrease in angiogenesis is indicated and as a factor in the wound healing process.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 176

The translation product of this gene shares sequence homology with MAT8 (mouse) which is thought to be important in regulating chloride conductance in cells (particularly in the breast) by modulating the response mediated by cAMP and protein kinase C to extracellular signals.

This gene is expressed primarily in amniotic cells and hematopoeitic cells including macrophages, Neutrophils, T cells, TNF induced aortic endothelium and to a lesser extent in testes, TNF induced epithelial cells, and smooth muscle.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, inflammatory responses mediated by T cells, macrophages, and/or neutrophils particularly those involving TNF, and also cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEO ID NO: 409 as residues: Thr-19 to Ala-33, Leu-54 to Asp-82, Pro-89 to Ala-97, Pro-100 to Lys-125, Ser-127 to Phe-135, Gly-164 to Leu-169, Cys-173 to Arg-178.

The tissue distribution and homology to mat-8 indicates that polynucleotides and polypeptides corresponding to this gene are useful for modifying inflammatory

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responses to cytokines such as TNF and thus modifying the duration and/or severity of inflammation. Polynucleotides and polypeptides derived from this gene are thought to be useful in the diagnosis and treatment of cancer.

# 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 177

This gene is expressed primarily in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, vascular restenosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating diseases associated with vascular response to injury such as vascular restenosis following angioplasty..

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 178

One embodiment of the claimed invention comprises:

- 25 MRPDWKAGAGPGGPPQKPAPSSQRKPPARPSAAAAAIAVAAAEEERRLRQRN RLRLEEDKPAVERCLEELVFGDVENDEDALLRRLRGPRVQEHEDSGDSEVENEA KGNFPPQKKPVWVDEEDEDEEMVDMMNNRFRKDMMKNASESKLSKDNLKK RLKEEFQHAMGGVPAWAETTKRKTSSDDESEEDEDDLLQRTGNFISTSTSLPRG ILKMKNCQHANAERPTVARISICAVPSRCTDCDGCWD (SEQ ID NO:737); or
- 30 CLEELVFGDVENDEDALLRRLRGPRVQEHEDSGDSEVENEAKGNFPPQKKPV
   WVDEEDEDEEMVDMMNNRFRKDMMKNASESKLSKDNLKKRLKEEFQHAMG
   GVPAWAETTKRKTSSDDESEEDEDDLLQRTGNFISTSTSLPRGILKMKNCQHA
   NAERPTVARISICAVPSRCTDCDGC (SEQ ID NO: 738). LKEKIVRSFEVSPDGS
   FLLINGIAGYLHLLAMKTKELIGSMKINGRVAASTFSSDSKKVYASSGDGEVYV

   35 WDVNSRKCLNRFVDEGSLYGLSIATSRNGQYVACGSNCGVVNIYNQDSCLOE
- TNPKPIKAIMNLVTGVTSLTFNPTTEILAIASEKMKEAVRLVHLPSCTVFSNFPVI KNKNISHVHTMDFSPRSGYFALGNEKGKALMYRLHHYSDF (SEQ ID NO:739);

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and/or KINGRVAASTFSSDSKKVYASSGDGEVYVWDVNSRKCLNRFVDEGSL YGLSIATSRNGQYVACGSNCGVVNIYNQDSCLQETNPKPIKAIMNLVTGVTSLT FNPTTEILAIASEKMKEAVRLVHLPSCTVFSNFPVIKNKNISHVHTMDFSPRSG YFALGNEKGKAL (SEQ ID NO:740).

This gene is expressed primarily in epidydimus and endometrial tumors and to a lesser extent in T cell lymphoma and cell lines derived from colon cancer.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, tumors of the reproductive organs including testis and endometrial cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 411 as residues: Ser-67 to Lys-72, Val-87 to Leu-93, Tyr-128 to Pro-141, Asp-204 to Gly-210.

The tissue distribution indicates that the protein products of this gene are useful for treating tumors of the endometrium or epithelial tumors of the reproductive system.

### 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 179

Preferred polypeptides encoded by this gene comprise the following amino acid sequence:

MRILQLILLALATGLVGGETRIIKGFECKLHSQPWQAALFEKTRLLCGATLIAPR WLLTAAHCLKPRYIVHLGQHNLQKEEGCEQTRTATESFPHPGFNNSLPNKDH RNDIMLVKMASPVSITWAVRPLTLSSRCVTAGTSCSFPAGAARPDPSYACLTPC DAPTSPSLSTRSVRTPTPATSQTPWCVPACRKGARTPARVTPGALWSVTSLFKA LSPGARIRVRSPESLVSTRKSANMWTGSRRR (SEQ ID NO:741); ETRIIKGFEC KLHSQPWQAALFEKTRLLCGATLIAPRWLLTAAHCLKPRYIVHLGQHNLQKEE GCEQTRTATESFPHPGFNNSLPNKDHRNDIMLVKMASPVSITWAVRPLTLSSR CVTAGTSCSFPAGAARPDPSYACLTPCDAPTSPSLSTRSVRTPTPATSQTPWCVP ACRKGARTPARVTPGALWSVTSLFKALSPGARIRVRSPESLVSTRKSANMWTG

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SRRR (SEQ ID NO:742); or CKLHSQPWQAALFEKTRLLCGATLIAPRWLLT AAHCLKPRYIVHLGQHNLQKEEGCEQTRTATESFPHPGFNS (SEQ ID NO:743). The translation product of this gene shares sequence homology with neuropsin a novel serine protease which is thought to be important in modulating

with neuropsin a novel serine protease which is thought to be important in modulating extracellular signaling pathways in the brain. Owing to the structural similarity to other serine proteases the protein products of this gene are expected to have serine protease activity which may be assayed by methods known in the art and described elsewhere herein.

This gene is expressed primarily in endometrial tumor and to a lesser extent in colon cancer, benign hypertrophic prostate, and thymus.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers of the endometrium or colon and benign hypertrophy of the prostate. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the urogenital or reproductive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 412 as residues: Gly-12 to Ser-22, Pro-34 to Ser-53.

The tissue distribution and homology to serine proteases indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosing or treating hyperproliferative disorders such as cancer of the endometrium or colon and hyperplasia of the prostate.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 180

Preferred polypeptide encoded by this gene comprise the following amino acid sequence: VLQGRYFSPILEMRRLRPEGXXNLPGGSRAQKEPRQDLTLVLWPHC PHFAMTRSYVPTKQCMVQGSFYCIFIFKGPVQNWC (SEQ ID NO:744).

Polynucleotides encoding such polypeptide are also provided.

This gene is expressed primarily in fetal brain

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, identifying and expanding stem cells in the CNS. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein products of this gene are useful for detecting and expanding stem cell populations in the (or of the) central nervous system.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 181

This gene is expressed primarily in early stage human brain and a stromal cell line.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, developmental abnormalities of the CNS. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 414 as residues: Gln-42 to Gln-47, Gln-54 to Pro-60.

The tissue distribution indicates that the protein products of this gene play a role in the development of the central nervous system. Therefore this gene and its products

are useful for diagnosing or treating developmental abnormalities of the central nervous system.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 182

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Preferred polypeptides encoded by this gene comprise the following amino acid sequence:

MPIIDQVNPELHDFMQSAEVGTIFALSWLITWFGHVLSDFRHVVRLYDF FLACHPLMPIYFAAVIVLYREQEVLDCDCDMASVHHLLSQIPQDLPYETLISRXE TFLFSFPHPNLLGRPLPNSKLRGRQPLLSKTLSWHQPSRGLIWCCGSGXRGLL RPEDRTKDVLTKPRTNRFVKLAVMGLTVALGAAALAVVKSALEWAPKFQLQL FP (SEQ ID NO:745); or CPEFFIPATLPCPFVFAFTSEASSRAYLTQRGPGGLAQ NLMPLPVGFWMGSLPPPWCWRKWVSEACSCFC (SEQ ID NO:746) These polypeptides are structurally similar to various TGF-beta family members. Thus, this polypeptide is expected to have a variety of activities in the modulation of cell growth and proliferation.

This gene is expressed primarily in osteoclastoma, microvascular endothelium, and bone marrow derived cell lines.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, hematological diseases particularly involving aberrant proliferation of stem cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 415 as residues: Ser-33 to Ala-39.

The tissue distribution indicates that the protein products of this gene is useful for treating disorders of the progenitors of the immune system. Applications include in vivo expansion of progenitor cells, ex vivo expansion of progenitor cells, or the treatment of tumors of the circulatory system, such as lymphomas.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 183

This gene maps to chromosome 17 and therefore, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 17. In specific embodiments, polypeptides of the invention comprise the sequence:

- 5 GFGSVSAAGRRSGGTWQPVQ (SEQ ID NO:747); PGGLAVGSRWWSRSLT (SEQ ID NO:748); LEPSRQRRPRRRGGTSRPETDQRAKCWRQL (SEQ ID NO:749); and/or VCLRCQNRMEN (SEQ ID NO:750). In further specific embodiments, polypeptides of the invention comprise the sequence: MAACTARRPGR GQPLVVPVADXGPVAKAALCAAXAGAFSPASTTTTRRHLSSRNRPEGKVLETV
- 10 GVFEVPKQNGKYETGQLFLHSIFGYRGVVLFPWQARLXDRDVASAAPEKAEN PAGHGSKEVKGKTHTYYQVLIDARDCPHISQRSQTEAVTFLANHDDSRALYAIP GLDYVSHEDILPYTSTDQVPIQHELFERFLLYDQTKAPPFVARETLRAWQEKNH PWLELSDVHRETTENIRVTVIPFYMGMREAQNSHVYWWRYCIRLENLDSDVVQ LRERHWRIFSLSGTLETVRGRGVVGREPVLSKEQPAFQYSSHVSLQASSGHMW
- 15 GTFRFERPDGSHFDVRIPPFSLESNKDEKTPPSGLHW (SEQ ID NO:751); MAACTARRPGRGQPLVVPVADXGPVAKAALCAA (SEQ ID NO:752); VLETVGVFEVPKQNGKYETGQLFLHSIFGYRGVVL (SEQ ID NO:757); GLDYVSHEDILPYTST (SEQ ID NO:758); DVHRETTENIRVTVIPFYM (SEQ ID NO:759); WWRYCIRLENLDSDVVQLRER (SEQ ID NO:760); and/or PAFQYSS
- HVSLQASSGHMWGTFRFER (SEQ ID NO:761). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in gall bladder, prostate, and fetal brain, and to a lesser extent in a few tumor and fetal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as 25 reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, growth related disorders such as cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders 30 of the above tissues or cells, particularly of the prostate, gall bladder, and fetal brain, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the 35 disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of growth-related disorders, such cancers.

### 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 184

In specific embodiments, polypeptides of the invention comprise the sequence:SLCCPEGAEGC (SEQ ID NO:762) and/or QLKKTHYDRPCP (SEQ ID NO:763). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in stromal cell, tonsil, and glioblastoma and to a lesser extent in some other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune and inflammatory disorders and glioblastoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the stromal cells, tonsil, and glioblastoma expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Additionally, it is believed that the product of this gene regulates pancreatic cell differentiation into beta cells. Accordingly, polynucleotides and polypeptides of the invention are useful in the treatment of insulindependent diabetes mellitus and associated conditions e.g. pancreatic hypofunction and the prevention, as well as the treatment of undifferentiated type pancreatic cancers. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 417 as residues: Pro-27 to Ala-32.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune and inflammatory disorders and glioblastoma.

# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 185

This gene is expressed primarily in hepatocellular carcinoma and to a lesser extent in other tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, liver diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the liver, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 418 as residues: Gly-32 to Lys-39.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of liver diseases.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 186

This gene is expressed primarily in hippocampus and to a lesser extent in other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neutronal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hippocampus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal disorders.

## 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 187

This gene is expressed primarily in bone cancer and hippocampus and to a lesser extent in osteoclastoma and other tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, bone-related disorders and neuronal diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the bone, ostoeclast, and hippocampus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of bone-related disorders and neuronal diseases.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 188

This gene maps to chromosome 4 and therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 4.

This gene is expressed primarily in neuronal tissues such as hippocampus, spinal cord, and hypothalamus and to a lesser extent in a few other tissues such as ovary.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal disorders.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 189

This gene maps to chromosome 10, therefore, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 10.

This gene is expressed primarily in neuronal tissues and immune tissues, and to a lesser extent in a few other tissues such as skin tumor, lung etc.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neuronal and immune-related disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuronal and immune-related tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 422 as residues: Pro-19 to Asp-25.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neuronal and immune-related disorders.

#### 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 190

The translation product of this gene shares sequence homology with human N33, a gene located in a homozygously deleted region of human metastatic prostate cancer which is thought to be important in prevention of prostate cancer. In specific embodiments, polypeptides of the invention comprise the sequence:

- AQRKKEMVLSEKVSQLMEWTNKRPVIRMNGDKFRRLVKAPPRNYSVIVMFTA
   LQLHRQCVVCKQADEEFQILANSWRYSSAFTNRIFFAMVDFDEGSDVFQMLNM
   NSAPTFINFPAKGKPKRGDTYELQVRGFSAEQIARWIADRTDVNIRVIRPPNMA
   ARWRFWCVSVT (SEQ ID NO:765); MVVALLIVCDVPSAS (SEQ ID NO:766);
   AQRKKEMVLSEKVSQL (SEQ ID NO:767); MEWTNKRPVIRMNGDKF (SEQ

   ID:768); RRLVKAPPRNYSVIVMFTALQLHRQCVVCKQADEEFQILANSWRY
  - SSAFTNRIFFA (SEQ ID NO:769); MVDFDEGSDVFQMLNMNSAPTFINFPAK GKP (SEQ ID NO:770); KRGDTYELQVRGFSAEQIARWIADRTDVNIRVIRPPN

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(SEQ ID NO:771); and/or YAGPLMLGLLLAVIGGLVYLRRVIWNFSLIKLDGLLQL CVLCLL (SEQ ID NO:772). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in infant adrenal gland prostate cell line and to a lesser extent in a few other tissues like liver, smooth muscle etc.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, prostate cancer and endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate and adrenal gland, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 423 as residues: Pro-34 to Gly-43, Arg-113 to Pro-120.

The tissue distribution and homology to N33 indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment for prostate cancer and endocrine disorders.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 191

This gene is expressed primarily in T cell and to a lesser extent in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue

or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 424 as residues: Trp-3 to Phe-9.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 192

This gene maps to chromosome 6, therefore, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 6. Neural activity and neurotrophins induce synaptic remodeling in part by altering gene expression. This gene is believed to be a glycosylphoshatidylinositol-anchored protein encoded by a hippocampal gene and to possess neural activity. This molecule is believed to be expressed in postmitotic-differentiating neurons of the developing nervous system and neuronal structures associated with plasticity in the adult. Message of this gene is believed to be induced by neuronal activity and by the activity-regulated neurotrophins BDNF and NT-3. The product of this gene is believed to stimulate neurite outgrowth and arborization in primary embryonic hippocampal and cortical cultures and to act as a downstream effector of activity-induced neurite outgrowth. In specific embodiments, polypeptides of the invention comprise the sequence: DAVFKGFSDCLLKLGDS (SEQ ID NO:773); CQEGAKDMWDKLRKESKNLN (SEQ ID NO:774);

20 VLLVSLSAALATWLSF (SEQ ID NO:775); MGLKLNGRYISLILAVQIAYLVQAVR AAGKCDAVFKGFSDCLLKLGDS (SEQ ID NO:776); PAAWDDKTNIKTVCTYW EDFHSCTVTALTDCQEGAKDMWDKLRKESKNLNIQGSLFELCGSGNGAAGSL LPAFPVLLVSLSAALATWLSF (SEQ ID NO:777); and/or MGLKLNGRYISLILA VQIAYLVQAVRAAGKCDAVFKGFSDCLLKLGDSXXXXXPAAWDDKTNIKTVC TYWEDFHSCTVTALTDCQEGAKDMWDKLRKESKNLNIQGSLFELCGSGNGAA GSLLPAFPVLLVSLSAALATWLSF (SEQ ID NO:778). Polynucleotides encoding this polypeptide are also encompassed by the invention.

This gene is expressed primarily in human placenta, endometrial tumor and tissues of the central nervous system (CNS).

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, relating to reproductive disorders, cancers and neurological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and neurological disorders, expression of this gene at significantly higher

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or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 425 as residues: Asp-47 to Asp-63, His-75 to Tyr-80, Pro-83 to Tyr-89.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of reproductive disorders such as endometrial tumors. Expression of this gene in tissues of the CNS and its strong homology to Neuritin suggest that the protein product from this gene may also be used in the treatment and diagnosis of neurological disorders and in the regeneration of neural tissues, e.g., following injury.

## 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 193

The translation product of this gene shares sequence homology with tenascin which is thought to be important in development. The translation product of this gene is believed to be a ligand of the fibroblast growth factor family. FGF ligand activity is known in the art and can be assayed by methods known in the art and disclosed elsewhere herein.

This gene is expressed primarily in endometrial tumors, and other types of tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cancer tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 426 as residues: Gly-29 to Glu-34, Arg-71 to Arg-76, Thr-176 to Cys-182, Gly-184 to Glu-199.

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The tissue distribution and homology to tenascin indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancers.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 194 5

In specific embodiments, polypeptides of the invention comprise the sequence: MNSAAGFSHLDRRERVLKLGESFEKQPRCASTLC (SEQ ID NO:779). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in fetal human lung and neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, lung development and respiratory disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the respiratory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution in fetal lung and neutrophils indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of lung and immunity related diseases, for example, lung cancer, viral, fungal or bacterial infections (e.g. lesions caused by tuberculosis), inflammation (e.g. pneumonia), metabolic lesions etc.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 195

This gene is expressed primarily in breast lymph node.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immunal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at

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significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of immunal disorders.

### 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 196

This gene maps to chromosome 5 and accordingly, polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 5. The translation product of this gene shares sequence homology with human M-phase phosphoprotein 4 which is thought to be important in phosphorylation and signal transduction processes. In specific embodiments, polypeptides of the invention comprise the sequence: TIYPTEEELQAVQKIVSITERALKLVSD (SEQ ID NO:780); RALKGVLRV GVLAKGLLLRGDRNVNLVLLC (SEQ ID NO:781); ALAALRHAKWFQARAN GLQSCVIIIRILRDLCQRVPTWS (SEQ ID NO:782); GDALRRVFECISSGIIL (SEQ ID NO:783); LAFRQIHKVLGMDPLP (SEQ ID NO:784); and/or TIYPTEELLOAVO KIVSITERALKLVSDSLSEHEKNKNKEGDDKKEGGKDRALKGVLRVGVLAKG LLLRGDRNVNLVLLCSEKPSKTLLSRIAENLPKQLAVISPEKYDIKCAVSEAAIIL NSCVEPKMQVTITLTSPIIREENMREGDVTSGMVKDPPDVLDROKCLDALAALR HAKWFQARANGLQSCVIIIRILRDLCQRVPTWSDFPSWAMELLVEKAISSASSP OSPGDALRRVFECISSGIILKGSPGLLDPCEKDPFDTLATMTDQQREDITSSAQFA LRLLAFRQIHKVLGMDPLPQMSQRFNIHNNRKRRRDSDGVDGFEAEGKKDKK DYDNF (SEQ ID NO:785). Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in Human Hippocampus and to a lesser extent in Prostate, Human Frontal Cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, disorders related to reproductive system and nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system and nervous system, expression of this gene at significantly higher or lower

WO 98/54963 PCT/US98/11422

153

levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to human M-phase phosphoprotein 4 indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of reproductive and nervous system disorders.

# 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 197

In specific embodiments, polypeptides of the invention comprise the sequence: MGSQHSAAARPSSCRRKQEDDRDG (SEQ ID NO:786); LLAEREQEEAIAQFPYVEFTGRDSITCLTC (SEQ ID NO:787); and/or QGTGYIPTEQVNELVALIPHSDQRLRPQRTKQYV (SEQ ID NO:788).

15 Polynucleotides encoding these polypeptides are also encompassed by the invention.

This gene is expressed primarily in Human Primary Breast Cancer and to a lesser extent in Human Adult Spleen, Hodgkin's Lymphoma I, Salivary Gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, cancer and immunal disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cancer and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 430 as residues: Ser-126 to Gly-138.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of cancer and immunal disorders.

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 198

This gene is expressed primarily in monocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, blood cell disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of blood cell disorders.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 199

This gene is expressed primarily in Human Ovary and Synovia and to a lesser extent in Human 8 Week Whole Embryo.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, reproductive and developmental disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and developmental system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of reproductive and developmental disorders.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 200

This gene maps to chromosome 8 and therefore polynucleotides of the invention can be used in linkage analysis as a marker for chromosome 8. The translation product of this gene shares limited sequence homology with collagen proline rich domain.

This gene is expressed primarily in CNS.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, neurological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 433 as residues: Pro-35 to Asp-41.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neurological diseases.

## 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 201

Translation product of this gene shares homology with a mammalian histone H1a protein. One embodiment for this gene is the polypeptide fragments comprising the following amino acid sequence: ARLNVGRESLKREMLKSQGVKVSESPMGAR HSSWPEGAAFCKKVQGAQMQFPPRR (SEQ ID NO:789); ARLNVGRESLKR EML (SEQ ID NO:790); LKSQGVKVSESPMGARHSSW (SEQ ID NO:791); AFCKKVQGAQMQFPPRR (SEQ ID NO:792). An additional embodiment is the polynucleotide fragments encoding these polypeptide (See Accession No. pirlS24178) fragments.

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are

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not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders. Since the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in vital immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 202

This gene is expressed primarily in neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders. Since the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 203

This gene is expressed primarily in Neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, infectious disorders, immune disorders, and cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 436 as residues: Thr-31 to Lys-36.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of infectious disorders, immune disorders, and cancers. Since the gene is expressed in cells of lymphoid origin, the natural gene product may be involved in immune functions. Therefore it may be also used as an agent for immunological disorders including arthritis, asthma, immune deficiency diseases such as AIDS, and leukemia. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and/or immunotherapy targets for the above listed tumors and tissues.

# FEATURES OF PROTEIN ENCODED BY GENE NO: 204

This gene maps to chromosome 16 and therefore polynucleotides of the invention can be used in linkage analysis as markers for chromosome 16. The translation product of this gene shares sequence homology with lactate dehydrogenase which is thought to be important in lactate metabolism.

This gene is expressed primarily in human tonsils and to a lesser extent in Spleen, and Neutrophils.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, immune disorders, infectious disorders, and cancers. Similarly,

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polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune disorders, infectious disorders, and cancers, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 437 as residues: Gly-7 to Ser-12.

The tissue distribution and homology to lactate dehydrogenase gene indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of immune disorders, infectious disorders, and cancers.

# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 205

The translation product of this gene shares sequence homology with Gcap1 protein which is developmentally regulated in brain.

This gene is expressed primarily in placenta and endometrial tumor and to a lesser extent in several other tumors.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, vasculogenesis/angiogenesis and tumorigenesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system and tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Gcap1 protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorder or dysfunction of vascular system of tumorigenesis.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 206

In specific embodiments, polypeptides of the invention comprise the sequence MPYAQWLAENDRFEEAQKAFHKAGRQREA (SEQ ID NO:799); VQVLEQLTNNAVAESRFNDAAYYYWMLSMQCLDIAQD (SEQ ID NO:794); PAQKDTMLGKFYHFQRLAELYHGYHAIHRHTEDP (SEQ ID NO: 795); FSVHRPETLFNISRFLLHSLPKDTPSGISKVKILFT (SEQ ID NO:800); LAKQSKALGAYRLARHAYDKLRGLYIP (SEQ ID NO:796); ARFQKSIELG TLTIRAKPFHDSEELVPLCYRCSTNN (SEQ ID NO: 797); and/or PLLNNLGNVC INCRQPFIFSASSYDVLHLVEFYLEEGITDEEAISLIDLEVLRPKRDDRQLEICKQQ LPDSCG (SEQ ID NO:798). Polynucleotides encoding these polypeptides are also

This gene is expressed primarily in testes.

encompassed by the invention.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 15 biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, male reproductive and endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive and endocrine systems, 20 expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. 25

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of male reproductive and endocrine disorders.

## 30 FEATURES OF PROTEIN ENCODED BY GENE NO: 207

This gene is expressed in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions, which include, but are not limited to, lung diseases such as cystic fibrosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

of the above tissues or cells, particularly of the respiratory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO: 440 as residues: Tyr-49 to Cys-54.

The tissue distribution indicates that polynucleotides and polypeptides

corresponding to this gene are useful for detection and treatment of disorders associated with developing lungs particularly in premature infants where the lungs are the last tissues to develop. The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and intervention of lung tumors since the gene may be involved in the regulation of cell division,

particularly since it is expressed in fetal tissue. Protein, as well as, antibodies directed against the protein may show utility as a tumor marker and immunotherapy targets for the above listed tumors and tissues.

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97974	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	209080 05/29/97	ATCC Deposit Nr and Date
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1821	1094	704	1404	1515	812	872		Total NT Seq.
892	1	22	1	118	p-a-sa	<u> </u>		5' NT of Clone Seq.
1647	1094	704	1265	1507	812	872		5' NT 3' NT of of Clone Clone Seq. Seq.
56	32		92	302	41	74		5' NT of Start Codon
56	32	117	92	302	41	74		of AA First SEQ AA of ID Signal NO: Pep Y
266	265	264	445	263	262	261		AA Firs SEQ AA ID of NO: Sig Y Pep
_	<b>–</b>		<b>—</b>	_	<u> </u>	1		First AA of Sig Pep
26	34	5	19	24	30	18		Last AA of Sig Pep
27	35	19	20	25	31	19		First AA of Secreted Portion
28	53	127	415	362	43	28		Last AA of ORF

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39	38	37	. 36	35	35	34		Gene No.		
HBMSN25	HATEF60	HAGFB60	HADAE74	HWTBF59	HWTBF59	HTXGI75		cDNA Clone ID		
97974	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	209080 05/29/97	Nr and Date	ATCC Deposit	
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pSport1	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	,	Vector		
49	48	47	46	223	45	44		XO:	E SEO	TIM
1742	2432	840	2421	707	983	1024		NT Seq.	Total	
1165	1193	- Pre-ma	664	488	779	30		Seq.	of Clone	7
1742   1165   1742	2246	840	1587	707	983	1024		Seq.	of of Clone	ינ זיי
1207	1491	97	710	514	85			Start Codon	5' NT	
1207	1491	97	710	514	85	167		Signal Pep	First AA of	5' NT
272	271	270	269	446	268	267		フ	D SES	•
	<b>}</b>	1		1	<b>)</b>	-		Sig Pep		1
23	17	30		41	30	20		Sig Pep	AA of	
24	18	31		42	31	21		Secreted Portion	First AA of	
31	51	48	2	64	221	25		of ORF		

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45	4	43	42	4	40		Gene No.
HCESF40	HCEEC15	HCECA49	HMDAN54	HCE3J79	HCDAR68		cDNA Clone ID
97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	04/04/97 209080 05/29/97	ATCC Deposit Nr and Date
pBluescript	Uni-ZAP XR		Vector				
55	54	53	52	51	50		× D. D. SEO
990	948	1558	1856	1328	1487		Total NT Seq.
99	<b>.</b>	310	725	251	181		5' NT of Clone Seq.
990	948	1408	1853	1328	1455		5' NT 3' NT of of Clone Clone Seq. Seq.
193	9	393	928	525	325		5' NT of Start
193	9	393	928	525	325		5' NT of First AA of Signal Pep
278	277	276	275	274	273		ΥÖ. BÖ SEÖ ¥
		. <u></u>	<b>—</b>	<b>—</b>	_		First AA of Sig Pep
32	23		33		35		Last AA of Sig Pep
33	24		34		36		First AA of Secreted Portion
256	65		50	21	56		Last AA of ORF

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51	50	49	48	47	46	45	Gene No.
HCWBB42	HCUDC07	HCRAF32	HCNAP62	HCMSX86	HCFMV39	HCESF40	cDNA Clone ID
97975 04/04/97 209081	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97974 04/04/97 209080 05/29/97	97974 04/04/97 209080 05/29/97	ATCC Deposit Nr and Date
ZAP Express	ZAP Express	Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	pSport1	pBluescript	Vector
61	60	59	58	57	56	224	NT SEQ ID NO.
618	478	1215	814	1052	1603	1384	Total NT Seq.
1		257	<b>,</b>	5	1	99	5' NT of Clone Seq.
618	478	1215	558	786	1296	1384	5' NT 3' NT of of Clone Clone Seq. Seq.
212	147		93	12	96	193	5' NT of Start Codon
212	147	356	93	12	96	193	of AA of SEQ AA of ID Signal NO: Pep Y
284	283	282	281	280	279	447	A SEQ
_	<u> </u>	· _	<u>-</u>	<b></b>	-	-	First AA of Sig Pep
35	36	19	22	28	29	32	First Last AA AA of of of Sig Sig Pep Pep
36	37	20	23	29	30	33	First AA of Secreted Portion
74	69	20	42	32	102	205	Last AA of ORF

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58	57	56	55	54	33	52		Gene No.
HE9HU17	HE6EU50	HE2OF09	HE2GS36	HE2AY71	HE2AV74	HDTAB05		cDNA Clone ID
97975 04/04/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pCMVSport 2.0		Vector
68	67	66	65	64	63	62		SEQ NO: NO:
2483	1152	1866	774	588	780	751		Total NT Seq.
1577	117	1313	272	21	283	<b></b>		5' NT of Clone Seq.
2448	686	1866	774	588	780	·751		5' NT 3' NT of of Clone Clone Seq. Seq.
1620	237	1596	445	169		257		S' NT of Start Codon
1620	237	1596	445	169	433	257		of First AA of Signal Pep
291	290	289	288	287	286	285		SEQ NO: U
	-	<u> </u>	1		-	-		First AA of Sig Pep
	20					21		First Last AA AA of of Sig Sig Pep Pep
	21					22		First AA of Secreted Portion
14	34	=	37	16	16	32		Last AA of ORF

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65	2	63	62	61	60	59		Gene No.	
HFVHY45	HFGAB89	HFEBA88	HEMAE80	HELDY74	HEBBW11	HE9ND48		cDNA Clone ID	
97975	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	209081 05/29/97	ATCC Deposit Nr and Date	
pBluescript	Uni-ZAP XR		Vector						
75	74	73	72	71	70			XO:	
831	1069	785	996	932	865	536		Total NT Seq.	
_	196	464	1	<b></b>	647	_		5' NT of Clone Seq.	
831	1047	785	945	932	865	536		5' NT 3' NT of Of Clone Clone Seq. Seq.	
	295	356	12	201		83		5' NT of Start Codon	
89	295	356	12	201	388	83		of First AA of Signal Pep	15' NT
298	297	296	295	294	293	292		AA SEQ ID NO:	
ഥ	<b>—</b>	<b></b>	<b></b>	-	-	-		Fep	
30	32	29	24	17	30	36		Last AA of Sig Pep	
31	ü	30	25	18	31	37		First AA of Secreted Portion	
76	3 <b>4</b>	57	136	33	135	43		Last AA of ORF	

71	70	69	68	67	66		Gene No.
HHGCN69	HHFHR32	ннғні59	ннгсг08	HGBBQ69	HGBAJ93		cDNA Clone ID
97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	04/04/97 209081 05/29/97	ATCC Deposit Nr and Date
Lambda ZAP II	Uni-ZAP XR		Vector				
81	80	79	78	77	76		X DE NT
1440	1378	661	1133	1274	590		Total NT Seq.
298	1	I	4	<b>—</b>	<b></b>		5' NT of Clone Seq.
1440	1378	661	1042	1273	590		5' NT 3' NT of of Clone Clone Seq. Seq.
532		192	175	105	233		5' NT of Start Codon
532	358	192	175	105	233		5' NT of First AA of Signal Pep
304	303	302	301	300	299		YÖ. BŞ SEQ SEQ
	-	<b>-</b>	-	-	<b>1–</b>		First AA of Sig Pep
23		29	23	24	38		Last AA of Sig Pep
24		30	24	25	39		First AA of Secreted Portion
34	13	112	30	43	94		Last AA of ORF

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82	82	80	79	78	77	76	75	74	73	72	Gene No.
HNGBT31	HNFJH45	HNFAE54	HMSKS35	HMEJE31	HKMNC43	HKIXL73	HJPAV06	HHSEG23	HHPFD63	HHGDO13	cDNA Clone ID
97976 04/04/97	97975 04/04/97 209081 05/29/97	97975 04/04/97 209081 05/29/97	ATCC Deposit Nr and Date								
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	pBluescript	pBluescript	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	Vector
92	91	90	89	88	87	98	85	84	83	82	XO:
639	575	1533	1102	655	908	1036	684	573	1706	1381	Total NT Seq.
		665	1	. 1	1	591	199	1	182	766	5' NT of Clone Seq.
639	575	1518	1102	655	908	1036	684	573	1644	1371	5' NT 3' NT of of Clone Clone Seq. Seq.
224	275	347	228	165	139	690	323	160	257	993	5' NT of Start Codon
224	275	347	228	165	139	690	323	160	257	993	5' NT of First AA of Signal Pep
315	314	313	312	311	310	309	308	307	306	305	AA SEQ ID NO:
_	-			-	-	-	-	-			First AA of Sig Pep
28	30	26	26	33	81	32	27	18	24	23	Last AA of Sig Pep
29	31	27	27	34	19	33	28	19	25	24	First AA of Secreted Portion
104	67	293	49	4	801	114	33	71	81	34	Last AA of ORF

91	90	89	88	8/	86	85	84	83	Gene No.	· · · · · ·	
HPCAL49	HPBCU51	HOSDI92	HOSBZ55	HOGAR52	HNHFL57	HNHDW42	HNGJG84	HNGIN60	cDNA Clone ID		
97977 04/04/97 209082	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	9/9// 04/04/97 209082 05/29/97	97976 04/04/97	97976 04/04/97	97976 04/04/97	97976 04/04/97	Nr and Date	ATCC Deposit	
Uni-ZAP XR	pBluescript SK-	Uni-ZAP XR	Uni-ZAP XR	pCMVSport 2.0	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Vector		
101	100	99	98	9/	96	95	94	93	×Ö	SEQ	T.
784	599	1935	1416	1985	844	426	526	744	NT Seq.	Total	
1	-	141	69	453		1	1	1	Seq.	of of Clone	Zi NI
784	599	772	1416	1985	844	426	526	744	Seq.	of of Of	اء الا
·	86		246	533	98	168	268	225	Start Codor	5' NT	
280	86	274	246	533	98	168,	268	225	Signal Pep	First AA of	of NT
324	323	322	321	320	319	318	317	316	ΥÖ	SEQ	AA
	<b>}</b>	1	P	-	-	-	1	1	Sig Pep	of ≱ }	Firet
81	27	20	32	1/	25	28	29		Sig Pep		1 281
19	28	21	33	18	26	29	30	44	Secreted Portion	First AA	
43	119	58	54	285	61	71	38		of ORF		

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97	96	95	95	94	93	92		Gene No.
HRGBR28	HRDFB85	HPWAN23	HPWAN23	HPMBQ32	НРНАС83	HPFCR13		cDNA Clone ID
97977 04/04/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector
107	106	226	105	104	103	102		NT SEQ ID NO:
107   1167	1705	2057	2066	1351	2218	1035		Total NT Seq.
611	23	1	51	_	840	602		5' NT of Clone Seq.
1167	1697	1954	2052	1351	2182	1035		5' NT 3' NT of of Clone Clone Seq. Seq.
53	233	220	270	18	1035	859		5' NT of Start Codon
53	233	220	270	81	1035	859		of of First AA of Signal Pep
330	329	449	328	327	326	325		Y. DSS A
	-	<b></b>	1	. –	1			First AA of Sig Pep
	21	29	29	23	17	32		Last AA of Sig Pep
2	22	30	30	. 24	81	33		First AA of Secreted Portion
263	201	315	537	86	17	58		Last AA of ORF

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102	101	100	100	99	98	98		Gene No.	
HTEFU09	HSXCS62	HSXBT86	HE8EU04	HSPAH56	HSKGN81	HSKGN81	•	cDNA Clone ID	
97977 04/04/97 209082	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209746 04/07/98	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209082 05/29/97	Deposit Nr and Date	
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pSport1	pBluescript	pBluescript		Vector	
112	111	228	110	109	227	108		× Ö E X	TN
2198	2249	2143	2632	611	2084	1907		Total NT Seq.	
228	1	53	294		335	151		of Clone Seq.	LŇ ,S
2158	1953	1096	2632	576	2084	1432		of of Clone Seq. Seq.	TŇ .6 LŇ .5
400	90	235	337	229	537	353		of Start Codon	
400	90	235	° 337	229	537	353		First AA of Signal Pep	
335	334	451	333	332	450	331		Y N E N	<b>A</b>
-	<b>)</b>	ь	_	H	-	<b></b>			First
	<del>-</del>		25	25	19	23		of Sig Pep	Last
	19		26	26	20	24		First AA of Secreted Portion	
23	199	9	333	47	.23	260		Last AA of ORF	

109	108	107	106	105	104	103		Gene No.
HTSHE40	HTSGM54	HTPCN79	НТОЕҮ16	HTGEW91	HTGEP89	нтекм35		cDNA Clone ID
97977 04/04/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	05/29/97	ATCC Deposit Nr and Date
pBluescript	pBluescript	Uni-ZAP XR		Vector				
119	118	117	116	115	114	113		SEQ NO:
119 1101	1133	503	1965	3684	703	1043		Total NT Seq.
118	316	1	127	526	_	40		5' NT of Clone Seq.
956	1069	503	1915	1338	703	1043		5' NT 3' NT of of Clone Clone Seq. Seq.
218	• .		202	584	285	320		5' NT of Start Codon
218	423	1	202	584	285	320		of AA of SEQ AA of ID Signal NO: Pep Y
342	341	340	339	338	337	336		AA SEQ ID NO: Y
-	1	1	1	1	1	1		First AA of Sig Pep
31	12	7	27	24	29	20		Last AA of Sig Pep
32	13	8	28	25	30	21		First AA of Secreted Portion
89	84	70	38	37	94	142		Last AA of ORF

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116	115	114	113	112		110		Gene No.
HE6EL90	HDTAW95	HCEVR60	HCE3Q10	HUKFC71	HTWBY29	HTWAF58		cDNA Clone ID
209007	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209082 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	pCMVSport 2.0	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	pSport1	Lambda ZAP II		Vector
126	125	124	123	122	121	120		NO: BO
1517	1288	1390	1542	994	2635	282		Total NT Seq.
_	412	82	<b>—</b>		1593			5' NT of Clone Seq.
1452	1288	1390	1542	932	2489	282		5' NT 3' NT of of Clone Clone Seq. Seq.
243	571	127	143		1654	137		5' No of Start Codo
243	571	127	143	272	1654	137	•	of of First AA of Signal Pep
349	348	347	346	345	344	343		YÖ. ⊞Ö SEÖ A
			<b>p</b>	-	-	-		AA First SEQ AA ID of NO: Sig Y Pep
		32	25	15	25	25		Last AA of Sig Pep
		33	26	. 16	26	26		First AA of Secreted Portion
9	16	153	63	221	55	48		Last AA of ORF

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122	121	120	119	18	117		Gene No.
HLTER03	HIBED17	ннРТD20	HFXBW82	HERAH36	HELBU29		cDNA Clone ID
209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	04/28/97 209083 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Other	Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	Uni-ZAP XR		Vector
132	131	130	129	128	127		XO BO
990	1950	472	1275	300	1073		Total NT Seq.
1	284	51		155	198		5' NT of Clone Seq.
990	1927	472	1275	300	1073		5' NT 3' NT of Of Clone Clone Seq. Seq.
78	395	·	56	202			5' NT of Start Codon
78	395	243	56	202	776		of First AA of Signal Pep
355	354	353	352	351	350		AA SEQ ID NO:
-	-	<b></b>		_	1		First AA of Sig Pep
22	72		23				Last AA of Sig Pep
23	73		24				First AA of Secreted Portion
34	245	32	61	17	13		Last AA of ORF

129	128	127	126	125	124	123	Gene No.
H6EAA53	HUKCO64	HSUBW09	HRGBR18	HPWAZ95	НРМСЈ92	HOABL56	cDNA Clone ID
209007 04/28/97 209083	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	Vector				
139	138	137	136	135	134	133	SEQ NO:
643	1777	1021	582	323	705	1720	Total NT Seg.
303	439	1	1		28	565	5' NT of Clone Seq.
643	1777	1021	582	323	705	1720	5' NT 3' NT of of Clone Clone Seq. Seq.
		153	·	88	106	660	5' NT of Start Codon
313	521	153	16	88	106	660	5' NT of First AA of Signal Pep
362	361	360	359	358	357	356	YO. DEQ
-	•	ь	H	<b>—</b>	<b>—</b>	1	First AA of Sig Pep
7		32	17	27	28	18	Last AA of Sig Pep
8			18	28	29	19	First AA of Secreted Portion
31	2	56	30	78	98	21	Last AA of ORF

135	134	134	133	132	- <del>-</del> 3	130		Gene No.	
HBMTD81	HBGCB91	HAIBP89	HALSQ59	HALSK07	HAGAO39	HAGAIII		cDNA Clone ID	
209008 04/28/97 209084 05/29/97	209007 04/28/97 209083 05/29/97	unknown 05/18/98	209007 04/28/97 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	209007 04/28/97 209083 05/29/97	05/29/97	ATCC Deposit Nr and Date	
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
145	229	144	143	142	141	140		SEQ ID NO:	Z
1082	1025	2243	300	1468	721	1220		Total NT Seq.	
163	409	173	4	125		. <b>L</b>		of Clone Seq.	LN 15
1082	1025	2243	300	1468	721	1220		of of Clone Seq. Seq. Seq.	יג ער
357	624	311	101	210				5' NT of Start Codon	,
357	624	311	101	210	415	127		First SEQ AA of ID Signal NO: Pep Y	S' NT
368	452	367	366	365	364	363		YOU SEO	
	<b> </b>	1	1	1	1	-		of Sig Pep	Firet
	20	27	22	29		16		of Sig Pep	T get
	21	28	23	30	•	17		First AA of Secreted Portion	
30	25	317	66	33	14	27		Last AA of ORF	

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142	141	140	139	138	137	136	Gene No.
HFCEB37	HE8EY43	HE2GT20	HCWHZ24	HCQAI40	HFKFJ07	HBXGK12	,cDNA Clone ID
209008 04/28/97 209084	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209010 04/28/97 209085 05/29/97	209008 04/28/97 209084 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	ZAP Express	Lambda ZAP II	Uni-ZAP XR	ZAP Express	Vector
152	151	150	149	148	147	146	NT SEQ ID NO:
802	2399	2890 `	1405	734	1183	4313	Total NT Seq.
352	1811	2890 1178	1	_	<b></b>	1153	5' NT of Clone Seq.
802	2399	2890	1405	734	1183	4313	5' NT 3' NT of of Clone Clone Seq. Seq.
	1265	1178	108	285	149	1313	5' NT of Start Codon
487	1265	1178	108	285	149	1313	5' NT of First AA of Signal Pep
375	374	373	372	371	370	369	Y. DEQ
-	-		<b></b>	-	-		First AA of Sig Pep
	30	31	34		41	18	Last AA of Sig Pep
	31	32	35		.42	. 19	First AA of Secreted Portion
10	34	39	63	19	254	42	Last AA of ORF

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149	148	147	146	145	144	143		Gene No.
HLMMU/6	HKLAB16	HUSIT49	НЈААU36	HHGBR15	HGLAM46	HFTCT67		cDNA Clone ID
209008	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	05/29/97	ATCC Deposit Nr and Date
Lambda ZAP II	Lambda ZAP II	pSport1	pBluescript SK-	Lambda ZAP II	Uni-ZAP XR	Uni-ZAP XR		Vector
159	158	157	156	155	154	153		× Ö. BÖ
1687	1625	157 2127	1251	642	2388	461		Total NT Seq.
1307	817	247	583	322	818	24		5' NT of Clone Seq.
1687	1625	2127	1251	642	2388	461		5' NT 3' NT of of Clone Clone Seq. Seq.
1296	1012	383		400	648	145		5' NT of Start
1296	1012	3 <u>8</u> 3	933	400	648	145		5' NT of First AA of Signal Pep
382	381	380	379	378	377	376		YÖ ⊞Ö A
_ –		<b></b>	<b>—</b>	-		Ь		First AA of Sig Pep
28	18	47	16			37	1	L Last AA Sig Pep
29		48	17			38		First AA of Secreted Portion
28	20	83	16	4	18	63		Last AA of ORF

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157	156	156	155	154	153	152	151	150		Gene No.	
H6EAE26	HSKCP69	HSKCP69	HPTRC15	HOECU83	HNHFQ63	HNHEJ88	HNH	HMS		Clor CD	
Æ26	CP69	CP69	CIS	CU83	FQ63	EJ88	HNHED86	HMSKQ35		cDNA Clone ID	
209009	209009 04/28/97	209009 04/28/97	209009 04/28/97	209009 04/28/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209008 04/28/97 209084 05/29/97	209084 05/29/97	Deposit Nr and Date	ATCC
9	)09 8/97	)09 8/97	)09 8/97	)09 3/97	)08 8/97 9/97	)08 8/97 384 )84 )/97	008 8/97 884 084 9/97	008 8/97 084 9/97	084 9/97	Deposit Nr and Date	.C
Uni-2	Uni-2	Uni-2	pBlı	Uni-2	Uni-	Uni-	Uni-	Uni-		,	
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	pBluescript	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
[ <del>S</del>		Ŕ	pt	ΚR	XR	XR	XR	XR			
167	230	166	165	164	163	162	161	160		X NO:	SEQ
882	1250	1251	2153	1400	753	519	770	1842		Total NT Seq.	
48	223	219	594	681	1	1		172			5' NT
882	1250	1120	2153	1400	753	519	770	1463		Clone Clone Seq. Seq.	5' NT 3' NT of
155	393				164	242	30	319		of Start Codon	5' NT
155	393		611	508	164	242	30	319		10	5' NT of First
390	453	389	388	387	386	385	384	383	_	Λö Nö T L	PAA SEQ
		-	<b></b>	-	_	-	-	-			First
33	32 ·			22	17	17	31	30		of Sig Pep	Last
34	33			23	18	18	32	31		of Secreted Portion	First AA
153	171		13	33	67	24	46	33			Last

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168	67	166	165	1 <u>6</u>	163	162	161	160	159	158		Gene No.	
HCFNF11	HCEZS40	HCEQA68	HCDDB78	HBMVP04	НВМТҮ28	HBHAD12	HAUAE83	HAICP19	HAGDQ47	HAGBX03		cDNA Clone ID	
209010	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209009 04/28/97	04/28/97	Nr and Date	ATCC							
pSport1	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
178	177	176	175	174	173	172	171	170	169	168		×ÖE	SEQ
1637	1502	1348	2379	888 888	1758	786	2003	170 1624	1307	1208		NT Seq.	
26	178	1	750	330	962	1	889	89	1	-			
1607	1502	1348	2379	862	1758	786	2003	1483	1307	1208		Clone Clone Seq. Seq.	5' NT 3' NT of
152	315	12	106		1184		1080	128	44	182		of Start Codon	TN 'S
152	315	12	901	546	1184	176	1080	128	44	182		AA of Signal Pep	
401	400	399	398	397	396	395	394	393	392	391		≺ÖB	S.
			-	-	-	. —		-	-			of Sig Pep	-
44		28	18		27	17		18	22			of Sig Pep	Last AA
45		29	19		28	18		19	23			of Secreted Portion	First AA
257	20	78	24	2	34	23	23	446	. 60	∞		ORF A	Last

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173	172	17.1	170	169	169		Gene No.
HE8MG65	HE2CT29	HDSAP81	HCUBL62	HCRBL20	HCRBL20		cDNA Clone ID
209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	04/28/97 209085 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	ZAP Express	Uni-ZAP XR	Uni-ZAP XR		Vector
183	182	181	180	231	179		NT SEQ ID NO:
2276	1128	968	519	1811	2911		Total NT Seq.
48	_	320	<b>—</b>	20	1103		5' NT of Clone Seq.
2276	1128	896	519	1811	2858		5' NT 3' NT of of Clone Clone Seq. Seq.
88	111	476	57	93	192		5' NT of Start Codon
88	111	476	. 57	93	192		of of First AA of Signal Pep
406	405	404	403	454	402		Y. DEQ SEQ
	<b></b> -	-	-	<u> </u>			First AA of Sig Pep
37	26	27	28	36	32		Last AA of Sig Pep
38	27	28	29	37	33		First AA of Secreted Portion
257	94	79	32	95	424		Last AA of ORF

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178	177	176	175	175	174	173	Gene No.
HETAR54	HEMDX17	HEMCV19	HEMAM41	HEMAM41	HE9FB42	HE8MG65	cDNA Clone ID
209010 04/28/97 209085	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	ATCC Deposit Nr and Date
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Vector
188	187	186	233	185	184	232	X NO: SEQ NO:
1848	654	941	1338	1337	2500	2271	Total NT Seq.
454	1	33	33	60	76	56	5' NT of Clone Seq.
1848	654	931	1327	1328	1693	2232	5' NT 3' NT of of Clone Clone Seq. Seq.
948	137	79	175	175	518	79	5' NT of Start Codon
948	137	79	175	175	518	79	5' NT of First AA of Signal Pep
411	410	409	456	408	407	455	AA SEC D NO: Y
	<b>}</b> -		_	· <b></b>	<b>—</b>	1	First AA of Sig Pep
14		23	32	39	-	43	First Last AA AA of of Sig Sig Pep Pep
15		24	33	40	2	44	First AA of Secreted Portion
232	13	178	91	190	623	170	Last AA of ORF

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187	186	185	184	183	182	181	180	179		Gene No.	
HHPSD37	HHPDW05	HHLBA89	HGLAM56	HGBF079	HFXHN68	HFKF140	HFGAB48	HETBX14		cDNA Clone ID	
209011 04/28/97	209011 04/28/97	209011 04/28/97	209011 04/28/97	209011 04/28/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	209010 04/28/97 209085 05/29/97	05/29/97	Deposit Nr and Date	ATCC
pBluescript	Uni-ZAP XR	pBluescript SK-	Uni-ZAP XR	Uni-ZAP XR	Lambda ZAP II	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR		Vector	
197	196	195	194	193	192	191	190	189		×ö. fl	SEO
1282	1443	1001	1098	1538	2118	1941	906	1146		Total NT Seq.	
66	1	1	68	259	777	120	156	157		Clone Seq.	of Of
1282	1443	1001	1098	1538	2118	1002	906	1146		Clone Clone Seq. Seq.	5' NT 3' NT
171	246	324		273	966	213	245			of Start Codon	TN 'S
171	246	324	185	273	966	213	245	74			5' NT of First
420	419	418	417	416	415	414	413	412		≺ÖÐ,	AA
	1	. –		_	<b>,</b>						First
19	.21	25	28	23	23	18	30	14		of Sig Pep	Last
20	22	26	29	24	24	19	31	15		of Secreted Portion	First AA
37	21	39	69	49	50	218	32	53			Last

			Γ		· -			·							
200	199	198	197	196	195	194	193	192	191	190	189	188	No.		
HNFAH08	HMSHQ24	HMSHM43	HLTDB65	нцтсү93	HLMIW92	HLHTC70	HLHSK94	нјрвв39	HJABZ65	HIASB53	HHSAK25	HHPSF70	Clone ID		
209011 04/28/97	Date	Deposit	ATCC												
Uni-ZAP XR	Lambda ZAP II	pBluescript	pBluescript	Uni-ZAP XR	pBluescript SK-	pBluescript	Uni-ZAP XR	pBluescript	Vector						
210	209	208	207	206	205	204	203	202	201	200	199	198	×Ċ	j B	SEO
2110	1779	872	1480	2465	721	1057	1974	1617	779	200   1707	1740	951	Seq.	Total	
592	16	1	1	886	1	229	1	188	1	401	1390	. 26	Seq.	Clone	5' N1
2110	1779	872	1480	2465	721	1057	1794	1605	779	1195	1740	951	Seq.	Clone Clone	5' NT 3' NT
119	148	35		1225	244	. 365	112	182	23	652	1534		Start Codon	of	
611	148	35	371	1225	244	365	112	182	23	652	1534	162	Signal Pep	AA of	5' NT of First
433	432	431	430	429	428	427	426	425	424	423	422	421	ΥĊ		AA
_	· -	<u> </u>	_	-	-	-	-		-	1	-	1	Sig	of.	First
18	24	18	15		25	23	26	28	26	26	19	16	Sig Pep		
19	25	19	16		26	24	27	29	27	27	20	17	Secreted Portion	of	First AA
191	36	36	143	4	46	22	379	91	68	126	31	34	$\overline{}$		Last

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207	206	205	204	203	202	201	Gene No.		
HCDE095	НРНАС88	HOSFM22	нинсм59	91ZVHNH	HNGBE45	010VOH	cDNA Clone ID		
209007 04/28/97 209083 05/29/97	97977 04/04/97 209082 05/29/97	97977 04/04/97 209082 05/29/97	209011 04/28/97	209011 04/28/97	209011 04/28/97	209011 04/28/97	Nr and Date	ATCC Deposit	
Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR	Uni-ZAP XR   214   1496	Uni-ZAP XR 213	Uni-ZAP XR 212 1551	Uni-ZAP XR	Vector		
217	216	215	214	213	212	211	×ö	E SEO	
999		1308	1496	997	1551	938	Seq.	Total	
608		501	_			-	Seq.	5' NT of Clone	
999			1132	997	1551	938	Seq.	S' NT 3' NT of Clone Clone	
273				202		107	Start Codon	Total Clone Clone of A	
273	549	809	165	202	114	107	Signal Pep	of AA First First SEQ AA AA of ID of	5' NT
440	439	438	437	436	435	434	۲. N	SEQ ID	•
-		-		_			Sig Pep	AA of	!
22	23		28	24	21	27	Sig Pep	AA of	•
23	24		29	25	22	28	Secreted Portion	AA First AA Last of of AA	
<b>54</b>	24		. 4	36	100	30	ORF of	Last AA	

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Table 1 summarizes the information corresponding to each "Gene No." described above. The nucleotide sequence identified as "NT SEQ ID NO:X" was assembled from partially homologous ("overlapping") sequences obtained from the "cDNA clone ID" identified in Table 1 and, in some cases, from additional related DNA clones. The overlapping sequences were assembled into a single contiguous sequence of high redundancy (usually three to five overlapping sequences at each nucleotide position), resulting in a final sequence identified as SEO ID NO:X.

The cDNA Clone ID was deposited on the date and given the corresponding deposit number listed in "ATCC Deposit No:Z and Date." Some of the deposits contain multiple different clones corresponding to the same gene. "Vector" refers to the type of vector contained in the cDNA Clone ID.

"Total NT Seq." refers to the total number of nucleotides in the contig identified by "Gene No." The deposited clone may contain all or most of these sequences, reflected by the nucleotide position indicated as "5' NT of Clone Seq." and the "3' NT of Clone Seq." of SEQ ID NO:X. The nucleotide position of SEQ ID NO:X of the putative start codon (methionine) is identified as "5' NT of Start Codon." Similarly, the nucleotide position of SEQ ID NO:X of the predicted signal sequence is identified as "5' NT of First AA of Signal Pep."

The translated amino acid sequence, beginning with the methionine, is identified as "AA SEQ ID NO:Y," although other reading frames can also be easily translated using known molecular biology techniques. The polypeptides produced by these alternative open reading frames are specifically contemplated by the present invention.

The first and last amino acid position of SEQ ID NO:Y of the predicted signal peptide is identified as "First AA of Sig Pep" and "Last AA of Sig Pep." The predicted first amino acid position of SEQ ID NO:Y of the secreted portion is identified as "Predicted First AA of Secreted Portion." Finally, the amino acid position of SEQ ID NO:Y of the last amino acid in the open reading frame is identified as "Last AA of ORF."

SEQ ID NO:X and the translated SEQ ID NO:Y are sufficiently accurate and otherwise suitable for a variety of uses well known in the art and described further below. For instance, SEQ ID NO:X is useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO:X or the cDNA contained in the deposited clone. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO:Y may be used to generate antibodies which bind specifically to the secreted proteins encoded by the cDNA clones identified in Table 1.

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Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

Accordingly, for those applications requiring precision in the nucleotide sequence or the amino acid sequence, the present invention provides not only the generated nucleotide sequence identified as SEQ ID NO:X and the predicted translated amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA containing a human cDNA of the invention deposited with the ATCC, as set forth in Table 1. The nucleotide sequence of each deposited clone can readily be determined by sequencing the deposited clone in accordance with known methods. The predicted amino acid sequence can then be verified from such deposits. Moreover, the amino acid sequence of the protein encoded by a particular clone can also be directly determined by peptide sequencing or by expressing the protein in a suitable host cell containing the deposited human cDNA, collecting the protein, and determining its sequence.

The present invention also relates to the genes corresponding to SEQ ID NO:X, SEQ ID NO:Y, or the deposited clone. The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

Also provided in the present invention are species homologs. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source for the desired homologue.

The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood in the art.

The polypeptides may be in the form of the secreted protein, including the mature form, or may be a part of a larger protein, such as a fusion protein (see below).

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It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple histidine residues, or an additional sequence for stability during recombinant production.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of a polypeptide, including the secreted polypeptide, can be substantially purified by the one-step method described in Smith and Johnson, Gene 67:31-40 (1988). Polypeptides of the invention also can be purified from natural or recombinant sources using antibodies of the invention raised against the secreted protein in methods which are well known in the art.

# Signal Sequences

Methods for predicting whether a protein has a signal sequence, as well as the cleavage point for that sequence, are available. For instance, the method of McGeoch, Virus Res. 3:271-286 (1985), uses the information from a short N-terminal charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje, Nucleic Acids Res. 14:4683-4690 (1986) uses the information from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 indicates the amino terminus of the secreted protein. The accuracy of predicting the cleavage points of known mammalian secretory proteins for each of these methods is in the range of 75-80%. (von Heinje, supra.) However, the two methods do not always produce the same predicted cleavage point(s) for a given protein.

In the present case, the deduced amino acid sequence of the secreted polypeptide was analyzed by a computer program called SignalP (Henrik Nielsen et al., Protein Engineering 10:1-6 (1997)), which predicts the cellular location of a protein based on the amino acid sequence. As part of this computational prediction of localization, the methods of McGeoch and von Heinje are incorporated. The analysis of the amino acid sequences of the secreted proteins described herein by this program provided the results shown in Table 1.

As one of ordinary skill would appreciate, however, cleavage sites sometimes vary from organism to organism and cannot be predicted with absolute certainty. Accordingly, the present invention provides secreted polypeptides having a sequence shown in SEQ ID NO:Y which have an N-terminus beginning within 5 residues (i.e., + or - 5 residues) of the predicted cleavage point. Similarly, it is also recognized that in some cases, cleavage of the signal sequence from a secreted protein is not entirely

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uniform, resulting in more than one secreted species. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

Moreover, the signal sequence identified by the above analysis may not necessarily predict the naturally occurring signal sequence. For example, the naturally occurring signal sequence may be further upstream from the predicted signal sequence. However, it is likely that the predicted signal sequence will be capable of directing the secreted protein to the ER. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

# 10 Polynucleotide and Polypeptide Variants

"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

By a polynucleotide having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the polypeptide. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the reference sequence. The query sequence may be an entire sequence shown in Table 1, the ORF (open reading frame), or any fragement specified as described herein.

As a practical matter, whether any particular nucleic acid molecule or polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleotide sequence of the presence invention can be determined conventionally using known computer programs. A preferred method for determing the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. (1990) 6:237-245). In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by converting U's to T's. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB alignment of DNA sequences to calculate percent identity are:

Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization

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Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty 0.05, Window Size=500 or the length of the subject nucleotide sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence because of 5' or 3' deletions, not because of internal deletions, a manual correction must be made to the results. This is because the FASTDB program does not account for 5' and 3' truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This corrected score is what is used for the purposes of the present invention. Only bases outside the 5' and 3' bases of the subject sequence, as displayed by the FASTDB alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

For example, a 90 base subject sequence is aligned to a 100 base query sequence to determine percent identity. The deletions occur at the 5' end of the subject sequence and therefore, the FASTDB alignment does not show a matched/alignement of the first 10 bases at 5' end. The 10 unpaired bases represent 10% of the sequence (number of bases at the 5' and 3' ends not matched/total number of bases in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 bases were perfectly matched the final percent identity would be 90%. In another example, a 90 base subject sequence is compared with a 100 base query sequence. This time the deletions are internal deletions so that there are no bases on the 5' or 3' of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only bases 5' and 3' of the subject sequence which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to made for the purposes of the present invention.

By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a query amino acid sequence of the present invention, it is intended that the amino acid sequence of the subject polypeptide is identical to the query sequence except that the subject polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the query amino acid sequence. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a query

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amino acid sequence, up to 5% of the amino acid residues in the subject sequence may be inserted, deleted, (indels) or substituted with another amino acid. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to, for instance, the amino acid sequences shown in Table 1 or to the amino acid sequence encoded by deposited DNA clone can be determined conventionally using known computer programs. A preferred method for determing the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. (1990) 6:237-245). In a sequence alignment the query and subject sequences are either both nucleotide sequences or both amino acid sequences. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB amino acid alignment are: Matrix=PAM 0, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Window Size=sequence length, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject amino acid sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence due to N- or Cterminal deletions, not because of internal deletions, a manual correction must be made to the results. This is becuase the FASTDB program does not account for N- and Cterminal truncations of the subject sequence when calculating global percent identity. 25 For subject sequences truncated at the N- and C-termini, relative to the the query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of 30 the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what is used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually adjusting the percent identity score. That is, 35 only query residue positions outside the farthest N- and C-terminal residues of the subject sequence.

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For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the Nterminus of the subject sequence and therefore, the FASTDB alignment does not show a matching/alignment of the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C- termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal deletions so there are no residues at the N- or Ctermini of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the FASTDB alignment, which are not matched/aligned with the query sequnce are manually corrected for. No other manual corrections are to made for the purposes of the present invention.

The variants may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as E. coli).

Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

Using known methods of protein engineering and recombinant DNA technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein without substantial loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988 (1993), reported variant KGF proteins having heparin binding activity even after

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deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).)

Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1a. They used random mutagenesis to generate over 3,500 individual IL-1a mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." (See, Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

Thus, the invention further includes polypeptide variants which show substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

WO 98/54963 PCT/US98/11422

The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham and Wells, Science 244:1081-1085 (1989).) The resulting mutant molecules can then be tested for biological activity.

As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

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## Polynucleotide and Polypeptide Fragments

In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in the deposited clone or shown in SEQ ID NO:X. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in the deposited clone or the nucleotide sequence shown in SEQ ID NO:X. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, 500, 600, 2000 nucleotides) are preferred.

Moreover, representative examples of polynucleotide fragments of the invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, 701-750, 751-800, 800-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150, 1151-1200, 1201-1250, 1251-1300, 1301-1350, 1351-1400, 1401-1450, 1451-1500, 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850, 1851-1900, 1901-1950, 1951-2000, or 2001 to the end of SEQ ID NO:X or the cDNA contained in the deposited clone. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has biological activity. More preferably, these polynucleotides can be used as probes or primers as discussed herein.

In the present invention, a "polypeptide fragment" refers to a short amino acid sequence contained in SEQ ID NO:Y or encoded by the cDNA contained in the deposited clone. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, or 161 to the end of the coding region. Moreover, polypeptide fragments can be about 20, 30, 40, 50, 60, 70, 80, 90, 100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the

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carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

Particularly, N-terminal deletions of the polypeptide of the present invention can be described by the general formula m-p, where p is the total number of amino acids in the polypeptide and m is an integer from 2 to (p-1), and where both of these integers (m & p) correspond to the position of the amino acid residue identified in SEQ ID NO:Y.

Moreover, C-terminal deletions of the polypeptide of the present invention can also be described by the general formula 1-n, where n is an integer from 2 to (p-1), and again where these integers (n & p) correspond to the position of the amino acid residue identified in SEQ ID NO:Y.

The invention also provides polypeptides having one or more amino acids deleted from both the amino and the carboxyl termini, which may be described generally as having residues m-n of SEQ ID NO:Y, where m and n are integers as described above.

20 Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alphahelix forming regions, beta-sheet and beta-sheet-forming regions, turn and turnforming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surfaceforming regions, substrate binding region, and high antigenic index regions. Polypeptide fragments of SEQ ID NO:Y falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

# Epitopes & Antibodies

35 In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred embodiment of the present invention relates to a polypeptide fragment comprising an

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epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA 81:3998- 4002 (1983).)

Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')2 fragments) which are capable of specifically binding to protein. Fab and F(ab')2 fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding than an intact antibody. (Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred, as well as the products of a FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

#### Fusion Proteins

Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the

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polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

Moreover, polypeptides of the present invention, including fragments, and specifically epitopes, can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life in vivo. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) can also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., J. Biochem. 270:3958-3964 (1995).)

Similarly, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. (EP-A 0232 262.) Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D.

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Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. Chem. 270:9459-9471 (1995).)

Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a peptide which facilitates purification of the fused polypeptide. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).)

Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

# 15 Vectors, Host Cells, and Protein Production

The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged in vitro using an appropriate packaging cell line and then transduced into host cells.

The polynucleotide insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination, and, in the transcribed region, a ribosome binding site for translation. The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin resistance for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance

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genes for culturing in E. coli and other bacteria. Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as E. coli, Streptomyces and Salmonella typhimurium cells; fungal cells, such as yeast cells; insect cells such as Drosophila S2 and Spodoptera Sf9 cells; animal cells such as CHO, COS, 293, and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan.

Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis et al., Basic Methods In Molecular Biology (1986). It is specifically contemplated that the polypeptides of the present invention may in fact be expressed by a host cell lacking a recombinant vector.

A polypeptide of this invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification.

Polypeptides of the present invention, and preferably the secreted form, can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any protein

after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient, depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

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### Uses of the Polynucleotides

Each of the polynucleotides identified herein can be used in numerous ways as reagents. The following description should be considered exemplary and utilizes known techniques.

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The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat polymorphisms), are presently available. Each polynucleotide of the present invention can be used as a chromosome marker.

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Briefly, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can be selected using computer analysis so that primers do not span more than one predicted exon in the genomic DNA. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the SEQ ID NO:X will yield an amplified fragment.

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Similarly, somatic hybrids provide a rapid method of PCR mapping the polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flow-sorted chromosomes, and preselection by hybridization to construct chromosome specific-cDNA libraries.

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Precise chromosomal location of the polynucleotides can also be achieved using fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides 2,000-4,000 bp are preferred. For a review of this technique, see Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York (1988).

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For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for marking multiple sites and/or multiple chromosomes). Preferred polynucleotides correspond to the noncoding regions of the cDNAs because the coding sequences are

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more likely conserved within gene families, thus increasing the chance of cross hybridization during chromosomal mapping.

Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library).) Assuming 1 megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential causative genes.

Thus, once coinheritance is established, differences in the polynucleotide and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using polynucleotides of the present invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker.

In addition to the foregoing, a polynucleotide can be used to control gene expression through triple helix formation or antisense DNA or RNA. Both methods rely on binding of the polynucleotide to DNA or RNA. For these techniques, preferred polynucleotides are usually 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991) ) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques are effective in model

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systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease.

Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell.

The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for RFLP.

The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992).) Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of

WO 98/54963

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unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers specific to particular tissue prepared from the sequences of the present invention. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for contamination.

In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response:

## Uses of the Polypeptides

Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

A polypeptide of the present invention can be used to assay protein levels in a biological sample using antibody-based techniques. For example, protein expression in tissues can be studied with classical immunohistological methods. (Jalkanen, M., et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell. Biol. 105:3087-3096 (1987).) Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (125I, 121I), carbon (14C), sulfur (35S), tritium (3H), indium (112In), and technetium (99mTc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

In addition to assaying secreted protein levels in a biological sample, proteins can also be detected in vivo by imaging. Antibody labels or markers for in vivo imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 131I, 112In, 99mTc), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously, or intraperitoneally) into the mammal. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. In vivo tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Tumor Imaging: The Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982).)

Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression of a polypeptide of the present invention in cells or body fluid of an individual; (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a disorder.

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Moreover, polypeptides of the present invention can be used to treat disease. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B), to inhibit the activity of a polypeptide (e.g., an oncogene), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth).

Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease. For example, administration of an antibody directed to a polypeptide of the present invention can bind and reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

At the very least, the polypeptides of the present invention can be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the following biological activities.

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### **Biological Activities**

The polynucleotides and polypeptides of the present invention can be used in assays to test for one or more biological activities. If these polynucleotides and polypeptides do exhibit activity in a particular assay, it is likely that these molecules may be involved in the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides could be used to treat the associated disease.

#### **Immune Activity**

A polypeptide or polynucleotide of the present invention may be useful in treating deficiencies or disorders of the immune system, by activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells from pluripotent stem cells. The etiology of these immune deficiencies or disorders may be genetic, somatic, such as cancer or some autoimmune disorders, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, a polynucleotide or polypeptide of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

A polynucleotide or polypeptide of the present invention may be useful in treating or detecting deficiencies or disorders of hematopoietic cells. A polypeptide or polynucleotide of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat those disorders associated with a decrease in certain (or many) types hematopoietic cells. Examples of immunologic deficiency syndromes include, but are not limited to: blood protein disorders (e.g. agammaglobulinemia, dysgammaglobulinemia), ataxia telangiectasia, common variable immunodeficiency, Digeorge Syndrome, HIV infection, HTLV-BLV infection, leukocyte adhesion deficiency syndrome, lymphopenia, phagocyte bactericidal dysfunction, severe combined immunodeficiency (SCIDs), Wiskott-Aldrich Disorder, anemia, thrombocytopenia, or hemoglobinuria.

Moreover, a polypeptide or polynucleotide of the present invention could also be used to modulate hemostatic (the stopping of bleeding) or thrombolytic activity (clot formation). For example, by increasing hemostatic or thrombolytic activity, a polynucleotide or polypeptide of the present invention could be used to treat blood coagulation disorders (e.g., afibrinogenemia, factor deficiencies), blood platelet disorders (e.g. thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, a polynucleotide or polypeptide of the present invention that can

WO 98/54963

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decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment of heart attacks (infarction), strokes, or scarring.

A polynucleotide or polypeptide of the present invention may also be useful in treating or detecting autoimmune disorders. Many autoimmune disorders result from inappropriate recognition of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing autoimmune disorders.

Examples of autoimmune disorders that can be treated or detected by the present invention include, but are not limited to: Addison's Disease, hemolytic anemia, antiphospholipid syndrome, rheumatoid arthritis, dermatitis, allergic encephalomyelitis, glomerulonephritis, Goodpasture's Syndrome, Graves' Disease, Multiple Sclerosis, Myasthenia Gravis, Neuritis, Ophthalmia, Bullous Pemphigoid, Pemphigus, Polyendocrinopathies, Purpura, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Thyroiditis, Systemic Lupus Erythematosus, Autoimmune Pulmonary Inflammation, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye disease.

Similarly, allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated by a polypeptide or polynucleotide of the present invention. Moreover, these molecules can be used to treat anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

A polynucleotide or polypeptide of the present invention may also be used to treat and/or prevent organ rejection or graft-versus-host disease (GVHD). Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. The administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing organ rejection or GVHD.

Similarly, a polypeptide or polynucleotide of the present invention may also be used to modulate inflammation. For example, the polypeptide or polynucleotide may inhibit the proliferation and differentiation of cells involved in an inflammatory response. These molecules can be used to treat inflammatory conditions, both chronic and acute conditions, including inflammation associated with infection (e.g., septic

shock, sepsis, or systemic inflammatory response syndrome (SIRS)), ischemia-reperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel disease, Crohn's disease, or resulting from over production of cytokines (e.g., TNF or IL-1.)

# **Hyperproliferative Disorders**

WO 98/54963

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A polypeptide or polynucleotide can be used to treat or detect hyperproliferative disorders, including neoplasms. A polypeptide or polynucleotide of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, a polypeptide or polynucleotide of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative disorders can be treated. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating hyperproliferative disorders, such as a chemotherapeutic agent.

Examples of hyperproliferative disorders that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but are not limited to neoplasms located in the: abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvic, skin, soft tissue, spleen, thoracic, and urogenital.

Similarly, other hyperproliferative disorders can also be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome, Waldenstron's Macroglobulinemia, Gaucher's Disease, histiocytosis, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

#### Infectious Disease

A polypeptide or polynucleotide of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases

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symptoms or diseases.

may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, the polypeptide or polynucleotide of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

Viruses are one example of an infectious agent that can cause disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of viruses, include, but are not limited to the following DNA and RNA viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus, Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza), Papovaviridae, Parvoviridae, Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g., Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis, bronchiollitis, encephalitis, eve infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), meningitis, opportunistic infections (e.g., AIDS), pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these

Similarly, bacterial or fungal agents that can cause disease or symptoms and that 25 can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following Gram-Negative and Gram-positive bacterial families and fungi: Actinomycetales (e.g., Corynebacterium, Mycobacterium, Norcardia), Aspergillosis, Bacillaceae (e.g., Anthrax, Clostridium), Bacteroidaceae, Blastomycosis, Bordetella, Borrelia, Brucellosis, Candidiasis, Campylobacter. 30 Coccidioidomycosis, Cryptococcosis, Dermatocycoses, Enterobacteriaceae (Klebsiella, Salmonella, Serratia, Yersinia), Erysipelothrix, Helicobacter, Legionellosis, Leptospirosis, Listeria, Mycoplasmatales, Neisseriaceae (e.g., Acinetobacter, Gonorrhea, Menigococcal), Pasteurellacea Infections (e.g., Actinobacillus, Heamophilus, Pasteurella), Pseudomonas, Rickettsiaceae, Chlamydiaceae, Syphilis, 35 and Staphylococcal. These bacterial or fungal families can cause the following diseases or symptoms, including, but not limited to: bacteremia, endocarditis, eye infections (conjunctivitis, tuberculosis, uveitis), gingivitis, opportunistic infections (e.g., AIDS

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related infections), paronychia, prosthesis-related infections, Reiter's Disease, respiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, Dysentery, Paratyphoid Fever, food poisoning, Typhoid, pneumonia, Gonorrhea, meningitis, Chlamydia, Syphilis, Diphtheria, Leprosy, Paratuberculosis, Tuberculosis, Lupus, Botulism, gangrene, tetanus, impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.

Moreover, parasitic agents causing disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following families: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis, Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis, Leishmaniasis, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas.

These parasites can cause a variety of diseases or symptoms, including, but not limited to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), Malaria, pregnancy complications, and toxoplasmosis. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.

Preferably, treatment using a polypeptide or polynucleotide of the present invention could either be by administering an effective amount of a polypeptide to the patient, or by removing cells from the patient, supplying the cells with a polynucleotide of the present invention, and returning the engineered cells to the patient (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease.

#### Regeneration

A polynucleotide or polypeptide of the present invention can be used to differentiate, proliferate, and attract cells, leading to the regeneration of tissues. (See, Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteocarthritis, periodontal disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion injury, or systemic cytokine damage.

Tissues that could be regenerated using the present invention include organs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal

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or cardiac), vascular (including vascular endothelium), nervous, hematopoietic, and skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs without or decreased scarring. Regeneration also may include angiogenesis.

Moreover, a polynucleotide or polypeptide of the present invention may increase regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. A polynucleotide or polypeptide of the present invention could also be used prophylactically in an effort to avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel syndrome, and other tendon or ligament defects. A further example of tissue regeneration of non-healing wounds includes pressure ulcers, ulcers associated with vascular insufficiency, surgical, and traumatic wounds.

Similarly, nerve and brain tissue could also be regenerated by using a polynucleotide or polypeptide of the present invention to proliferate and differentiate nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and stoke). Specifically, diseases associated with peripheral nerve injuries, peripheral neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome), could all be treated using the polynucleotide or polypeptide of the present invention.

#### **Chemotaxis**

A polynucleotide or polypeptide of the present invention may have chemotaxis activity. A chemotaxic molecule attracts or mobilizes cells (e.g., monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells) to a particular site in the body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

A polynucleotide or polypeptide of the present invention may increase chemotaxic activity of particular cells. These chemotactic molecules can then be used to treat inflammation, infection, hyperproliferative disorders, or any immune system disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotaxic molecules can be used to treat wounds and other trauma to tissues by attracting immune cells to the injured location. Chemotactic molecules of the present invention can also attract fibroblasts, which can be used to treat wounds.

WO 98/54963 PCT/US98/11422

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It is also contemplated that a polynucleotide or polypeptide of the present invention may inhibit chemotactic activity. These molecules could also be used to treat disorders. Thus, a polynucleotide or polypeptide of the present invention could be used as an inhibitor of chemotaxis.

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#### **Binding Activity**

A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules.

Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

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Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide, either as a secreted protein or on the cell membrane. Preferred cells include cells from mammals, yeast, Drosophila, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the molecule to observe binding, stimulation, or inhibition of activity of either the polypeptide or the molecule.

The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

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Alternatively, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a standard.

WO 98/54963

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Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptide from suitably manipulated cells or tissues.

Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

### **Other Activities**

A polypeptide or polynucleotide of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

A polypeptide or polynucleotide of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide or polynucleotide of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

A polypeptide or polynucleotide of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, caricadic rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity), hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

A polypeptide or polynucleotide of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional components.

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## Other Preferred Embodiments

Other preferred embodiments of the claimed invention include an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1.

Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Clone Sequence and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Start Codon and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Similarly preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the 5' Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 150 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

Further preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 500 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of SEQ ID NO:X beginning with the nucleotide at about the position of the 5' Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

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A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NO:X.

Also preferred is an isolated nucleic acid molecule which hybridizes under stringent hybridization conditions to a nucleic acid molecule, wherein said nucleic acid molecule which hybridizes does not hybridize under stringent hybridization conditions to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or of only T residues.

Also preferred is a composition of matter comprising a DNA molecule which comprises a human cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the material deposited with the American Type Culture Collection and given the ATCC Deposit Number shown in Table 1 for said cDNA Clone Identifier.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in the nucleotide sequence of a human cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the deposit given the ATCC Deposit Number shown in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of the complete open reading frame sequence encoded by said human cDNA clone.

Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by said human cDNA clone.

A further preferred embodiment is a method for detecting in a biological sample a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method

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comprises a step of comparing a nucleotide sequence of at least one nucleic acid molecule in said sample with a sequence selected from said group and determining whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95%

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identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1.

Also preferred is a polypeptide, wherein said sequence of contiguous amino acids is included in the amino acid sequence of SEQ ID NO:Y in the range of positions beginning with the residue at about the position of the First Amino Acid of the Secreted Portion and ending with the residue at about the Last Amino Acid of the Open Reading Frame as set forth for SEQ ID NO:Y in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a secreted portion of the secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

WO 98/54963

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Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the amino acid sequence of the secreted portion of the protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is an isolated antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method comprises a step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group and determining whether the sequence of said polypeptide molecule in said sample is at least 90% identical to said sequence of at least 10 contiguous amino acids.

Also preferred is the above method wherein said step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of polypeptides in said sample to an antibody which binds specifically to a polypeptide

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comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

Also preferred is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting polypeptide molecules in said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is the above method for identifying the species, tissue or cell type of a biological sample, which method comprises a step of detecting polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the above group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

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Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding a polypeptide wherein said polypeptide comprises an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said nucleotide sequence encoding a polypeptide has been optimized for expression of said polypeptide in a prokaryotic host.

Also preferred is an isolated nucleic acid molecule, wherein said polypeptide comprises an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is a method of making a recombinant vector comprising inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is the recombinant vector produced by this method. Also preferred is a method of making a recombinant host cell comprising introducing the vector into a host cell, as well as the recombinant host cell produced by this method.

Also preferred is a method of making an isolated polypeptide comprising culturing this recombinant host cell under conditions such that said polypeptide is expressed and recovering said polypeptide. Also preferred is this method of making an isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said polypeptide is a secreted portion of a human secreted protein comprising an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y beginning with the residue at the position of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y wherein Y is an integer set forth in Table 1 and said position of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y is defined in Table 1; and an amino acid sequence of a secreted portion of a protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The isolated polypeptide produced by this method is also preferred.

Also preferred is a method of treatment of an individual in need of an increased level of a secreted protein activity, which method comprises administering to such an individual a pharmaceutical composition comprising an amount of an isolated polypeptide, polynucleotide, or antibody of the claimed invention effective to increase the level of said protein activity in said individual.

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

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#### **Examples**

## Example 1: Isolation of a Selected cDNA Clone From the Deposited Sample

Each cDNA clone in a cited ATCC deposit is contained in a plasmid vector. Table 1 identifies the vectors used to construct the cDNA library from which each clone was isolated. In many cases, the vector used to construct the library is a phage vector from which a plasmid has been excised. The table immediately below correlates the related plasmid for each phage vector used in constructing the cDNA library. For example, where a particular clone is identified in Table 1 as being isolated in the vector "Lambda Zap," the corresponding deposited clone is in "pBluescript."

Vector Used to Construct Library Corresponding Deposited Plasmid Lambda Zap pBluescript (pBS) Uni-Zap XR pBluescript (pBS) Zap Express pBK 25 lafmid BA plafmid BA pSport1 pSport1 pCMVSport 2.0 pCMVSport 2.0 pCMVSport 3.0 pCMVSport 3.0 pCR<sup>®</sup>2.1 pCR<sup>®</sup>2.1

Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res. 16:7583-7600 (1988); Alting-Mees, M. A. and Short, J. M., Nucleic Acids Res. 17:9494 (1989)) and pBK (Alting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1

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Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS. The S and K refers to the orientation of the polylinker to the T7 and T3 primer sequences which flank the polylinker region ("S" is for SacI and "K" is for KpnI which are the first sites on each respective end of the linker). "+" or "-" refer to the orientation of the f1 origin of replication ("ori"), such that in one orientation, single stranded rescue initiated from the f1 ori generates sense strand DNA and in the other, antisense.

Vectors pSport1, pCMVSport 2.0 and pCMVSport 3.0, were obtained from Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gruber, C. E., et al., Focus 15:59 (1993).) Vector lafmid BA (Bento Soares, Columbia University, NY) contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1 Blue. Vector pCR®2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark, J. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9: (1991).) Preferably, a polynucleotide of the present invention does not comprise the phage vector sequences identified for the particular clone in Table 1, as well as the corresponding plasmid vector sequences designated above.

The deposited material in the sample assigned the ATCC Deposit Number cited in Table 1 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing the same ATCC Deposit Number contain at least a plasmid for each cDNA clone identified in Table 1. Typically, each ATCC deposit sample cited in Table 1 comprises a mixture of approximately equal amounts (by weight) of about 50 plasmid DNAs, each containing a different cDNA clone; but such a deposit sample may include plasmids for more or less than 50 cDNA clones, up to about 500 cDNA clones.

Two approaches can be used to isolate a particular clone from the deposited sample of plasmid DNAs cited for that clone in Table 1. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to SEQ ID NO:X.

Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with <sup>32</sup>P-γ-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).)

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The plasmid mixture is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1.93 to 1.104), or other techniques known to those of skill in the art.

Alternatively, two primers of 17-20 nucleotides derived from both ends of the SEQ ID NO:X (i.e., within the region of SEQ ID NO:X bounded by the 5' NT and the 3' NT of the clone defined in Table 1) are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 µl of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl<sub>2</sub>, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., Nucleic Acids Res. 21(7):1683-1684 (1993).)

Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.

This above method starts with total RNA isolated from the desired source, although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA which may interfere with the later RNA ligase step. The phosphatase should then be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.

This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene.

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## Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the cDNA sequence corresponding to SEQ ID NO:X., according to the method described in Example 1. (See also, Sambrook.)

## **Example 3: Tissue Distribution of Polypeptide**

Tissue distribution of mRNA expression of polynucleotides of the present invention is determined using protocols for Northern blot analysis, described by, among others, Sambrook et al. For example, a cDNA probe produced by the method described in Example 1 is labeled with P<sup>32</sup> using the rediprime<sup>™</sup> DNA labeling system (Amersham Life Science), according to manufacturer's instructions. After labeling, the probe is purified using CHROMA SPIN-100<sup>™</sup> column (Clontech Laboratories, Inc.), according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various human tissues for mRNA expression.

Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) (Clontech) are examined with the labeled probe using ExpressHyb™ hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are mounted and exposed to film at -70°C overnight, and the films developed according to standard procedures.

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#### Example 4: Chromosomal Mapping of the Polynucleotides

An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This primer set is then used in a polymerase chain reaction under the following set of conditions: 30 seconds, 95°C; 1 minute, 56°C; 1 minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions is analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

### Example 5: Bacterial Expression of a Polypeptide

A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and XbaI, at the 5' end of the primers in order to clone the amplified product into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (Amp<sup>r</sup>), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (Kan<sup>r</sup>). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis.

Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D. 600) of between 0.4 and 0.6. IPTG

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(Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression.

Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is removed by centrifugation, and the supernatant containing the polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., *supra*).

Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins are eluted by the addition of 250 mM immidazole. Immidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

In addition to the above expression vector, the present invention further includes an expression vector comprising phage operator and promoter elements operatively linked to a polynucleotide of the present invention, called pHE4a. (ATCC Accession Number 209645, deposited on February 25, 1998.) This vector contains: 1) a neomycinphosphotransferase gene as a selection marker, 2) an E. coli origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgarno sequence, and 6) the lactose operon repressor gene (lacIq). The origin of replication (oriC) is derived from pUC19 (LTI, Gaithersburg, MD). The promoter sequence and operator sequences are made synthetically.

DNA can be inserted into the pHEa by restricting the vector with NdeI and XbaI, BamHI, XhoI, or Asp718, running the restricted product on a gel, and isolating the larger fragment (the stuffer fragment should be about 310 base pairs). The DNA

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insert is generated according to the PCR protocol described in Example 1, using PCR primers having restriction sites for NdeI (5' primer) and XbaI, BamHI, XhoI, or Asp718 (3' primer). The PCR insert is gel purified and restricted with compatible enzymes. The insert and vector are ligated according to standard protocols.

The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

## Example 6: Purification of a Polypeptide from an Inclusion Body

The following alternative method can be used to purify a polypeptide expressed in *E coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells are then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the polypeptide containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with  $0.16\,\mu m$  membrane filter with appropriate surface area

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(e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A<sub>280</sub> monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

# Example 7: Cloning and Expression of a Polypeptide in a Baculovirus Expression System

In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide into a baculovirus to express a polypeptide. This expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak Drosophila promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

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Many other baculovirus vectors can be used in place of the vector above, such as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation, secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow et al., Virology 170:31-39 (1989).

Specifically, the cDNA sequence contained in the deposited clone, including the AUG initiation codon and the naturally associated leader sequence identified in Table 1, is amplified using the PCR protocol described in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the pA2 vector does not need a second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures," Texas Agricultural Experimental Station Bulletin No. 1555 (1987).

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("Geneclean" BIO 101 Inc., La Jolla, Ca.).

The fragment and the dephosphorylated plasmid are ligated together with T4 DNA ligase. E. coli HB101 or other suitable E. coli hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing.

Five μg of a plasmid containing the polynucleotide is co-transfected with 1.0 μg of a commercially available linearized baculovirus DNA ("BaculoGold<sup>TM</sup> baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One μg of BaculoGold<sup>TM</sup> virus DNA and 5 μg of the plasmid are mixed in a sterile well of a microtiter plate containing 50 μl of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10 μl Lipofectin plus 90 μl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm

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tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, *supra*. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 µl of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5  $\mu$ Ci of <sup>35</sup>S-methionine and 5  $\mu$ Ci <sup>35</sup>S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

## 30 Example 8: Expression of a Polypeptide in Mammalian Cells

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates

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the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLVI, HIVI and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSport 2.0, and pCMVSport 3.0. Mammalian host cells that could be used include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transfected cells.

The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing 20 cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Page, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991).) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); 25 Bebbington et al., Bio/Technology 10:169-175 (1992). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No.209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985).) Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the cloning of the gene of interest. The vectors also contain the 3' intron, the

.WO 98/54963 PCT/US98/11422

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polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

A polynucleotide of the present invention is amplified according to the protocol outlined in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the vector does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene is used for 20 transfection. Five µg of the expression plasmid pC6 is cotransfected with 0.5 µg of the plasmid pSVneo using lipofectin (Felgner et al., supra). The plasmid pSV2-neo contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are 25 trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of metothrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of 30 methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 µM, 2 µM, 5 µM, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 -200 µM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

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## **Example 9: Protein Fusions**

The polypeptides of the present invention are preferably fused to other proteins. These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding protein facilitates purification. (See Example 5; see also EP A 394,827; Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and albumin increases the halflife time in vivo. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an expression vector, preferably a mammalian expression vector.

For example, if pC4 (Accession No. 209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

If the naturally occurring signal sequence is used to produce the secreted protein, pC4 does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

### Human IgG Fc region:

GGGATCCGGAGCCCAAATCTTCTGACAAAACTCACACATGCCCACCGTGCC
CAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCCTCTTCCCCCCAAAACC
35 CAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTCACATGCGTGGTGGT
GGACGTAAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACG
GCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAAC

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AGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACTGGCTG
AATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAACCCCC
ATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGT
GTACACCCTGCCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTCAGCCT
GACCTGCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGTGGAGTGGGA
GAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGCTGG
ACTCCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCA
GGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGC
ACAACCACTACACGCAGAAGACCTCCCCTGTCTCCCGGGTAAATGAGTGC
GACGCCGCGACTCTAGAGGAT (SEQ ID NO:1)

## Example 10: Production of an Antibody from a Polypeptide

The antibodies of the present invention can be prepared by a variety of methods. (See, Current Protocols, Chapter 2.) For example, cells expressing a polypeptide of the present invention is administered to an animal to induce the production of sera containing polyclonal antibodies. In a preferred method, a preparation of the secreted protein is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

In the most preferred method, the antibodies of the present invention are monoclonal antibodies (or protein binding fragments thereof). Such monoclonal antibodies can be prepared using hybridoma technology. (Köhler et al., Nature 256:495 (1975); Köhler et al., Eur. J. Immunol. 6:511 (1976); Köhler et al., Eur. J. Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell Hybridomas, Elsevier, N.Y., pp. 563-681 (1981).) In general, such procedures involve immunizing an animal (preferably a mouse) with polypeptide or, more preferably, with a secreted polypeptide-expressing cell. Such cells may be cultured in any suitable tissue culture medium; however, it is preferable to culture cells in Earle's modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about 1,000 U/ml of penicillin, and about 100 μg/ml of streptomycin.

The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are selectively maintained in HAT medium, and then cloned by limiting dilution as

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described by Wands et al. (Gastroenterology 80:225-232 (1981).) The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the polypeptide.

Alternatively, additional antibodies capable of binding to the polypeptide can be produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the protein-specific antibody can be blocked by the polypeptide. Such antibodies comprise anti-idiotypic antibodies to the protein-specific antibody and can be used to immunize an animal to induce formation of further protein-specific antibodies.

It will be appreciated that Fab and F(ab')2 and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such fragments are typically produced by proteolytic cleavage, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). Alternatively, secreted protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry.

For in vivo use of antibodies in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art. (See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985).)

Example 11: Production Of Secreted Protein For High-Throughput

Screening Assays

The following protocol produces a supernatant containing a polypeptide to be tested. This supernatant can then be used in the Screening Assays described in Examples 13-20.

First, dilute Poly-D-Lysine (644 587 Boehringer-Mannheim) stock solution (1mg/ml in PBS) 1:20 in PBS (w/o calcium or magnesium 17-516F Biowhittaker) for a

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working solution of 50ug/ml. Add 200 ul of this solution to each well (24 well plates) and incubate at RT for 20 minutes. Be sure to distribute the solution over each well (note: a 12-channel pipetter may be used with tips on every other channel). Aspirate off the Poly-D-Lysine solution and rinse with 1ml PBS (Phosphate Buffered Saline). The PBS should remain in the well until just prior to plating the cells and plates may be poly-lysine coated in advance for up to two weeks.

Plate 293T cells (do not carry cells past P+20) at 2 x 10<sup>5</sup> cells/well in .5ml DMEM(Dulbecco's Modified Eagle Medium)(with 4.5 G/L glucose and L-glutamine (12-604F Biowhittaker))/10% heat inactivated FBS(14-503F Biowhittaker)/1x Penstrep(17-602E Biowhittaker). Let the cells grow overnight.

The next day, mix together in a sterile solution basin: 300 ul Lipofectamine (18324-012 Gibco/BRL) and 5ml Optimem I (31985070 Gibco/BRL)/96-well plate. With a small volume multi-channel pipetter, aliquot approximately 2ug of an expression vector containing a polynucleotide insert, produced by the methods described in Examples 8 or 9, into an appropriately labeled 96-well round bottom plate. With a multi-channel pipetter, add 50ul of the Lipofectamine/Optimem I mixture to each well. Pipette up and down gently to mix. Incubate at RT 15-45 minutes. After about 20 minutes, use a multi-channel pipetter to add 150ul Optimem I to each well. As a control, one plate of vector DNA lacking an insert should be transfected with each set of transfections.

Preferably, the transfection should be performed by tag-teaming the following tasks. By tag-teaming, hands on time is cut in half, and the cells do not spend too much time on PBS. First, person A aspirates off the media from four 24-well plates of cells, and then person B rinses each well with .5-1ml PBS. Person A then aspirates off PBS rinse, and person B, using a12-channel pipetter with tips on every other channel, adds the 200ul of DNA/Lipofectamine/Optimem I complex to the odd wells first, then to the even wells, to each row on the 24-well plates. Incubate at 37°C for 6 hours.

While cells are incubating, prepare appropriate media, either 1%BSA in DMEM with 1x penstrep, or CHO-5 media (116.6 mg/L of CaCl2 (anhyd); 0.00130 mg/L

CuSO<sub>4</sub>-5H<sub>2</sub>O; 0.050 mg/L of Fe(NO<sub>3</sub>)<sub>3</sub>-9H<sub>2</sub>O; 0.417 mg/L of FeSO<sub>4</sub>-7H<sub>2</sub>O; 311.80 mg/L of Kcl; 28.64 mg/L of MgCl<sub>2</sub>; 48.84 mg/L of MgSO<sub>4</sub>; 6995.50 mg/L of NaCl; 2400.0 mg/L of NaHCO<sub>3</sub>; 62.50 mg/L of NaH<sub>2</sub>PO<sub>4</sub>-H<sub>2</sub>O; 71.02 mg/L of Na<sub>2</sub>HPO4; .4320 mg/L of ZnSO<sub>4</sub>-7H<sub>2</sub>O; .002 mg/L of Arachidonic Acid; 1.022 mg/L of Cholesterol; .070 mg/L of DL-alpha-Tocopherol-Acetate; 0.0520 mg/L of Linoleic Acid; 0.010 mg/L of Linolenic Acid; 0.010 mg/L of Palmitic Acid; 0.010 mg/L of Palmitic Acid; 100 mg/L of

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Pluronic F-68; 0.010 mg/L of Stearic Acid; 2.20 mg/L of Tween 80; 4551 mg/L of D-Glucose; 130.85 mg/ml of L- Alanine; 147.50 mg/ml of L-Arginine-HCL; 7.50 mg/ml of L-Asparagine-H<sub>2</sub>0; 6.65 mg/ml of L-Aspartic Acid; 29.56 mg/ml of L-Cystine-2HCL-H,0; 31.29 mg/ml of L-Cystine-2HCL; 7.35 mg/ml of L-Glutamic Acid; 365.0 5 mg/ml of L-Glutamine; 18.75 mg/ml of Glycine; 52.48 mg/ml of L-Histidine-HCL-H<sub>2</sub>0; 106.97 mg/ml of L-Isoleucine; 111.45 mg/ml of L-Leucine; 163.75 mg/ml of L-Lysine HCL; 32.34 mg/ml of L-Methionine; 68.48 mg/ml of L-Phenylalainine; 40.0 mg/ml of L-Proline; 26.25 mg/ml of L-Serine; 101.05 mg/ml of L-Threonine; 19.22 mg/ml of L-Tryptophan; 91.79 mg/ml of L-Tryrosine-2Na-2H,0; 99.65 mg/ml of L-10 Valine; 0.0035 mg/L of Biotin; 3.24 mg/L of D-Ca Pantothenate; 11.78 mg/L of Choline Chloride; 4.65 mg/L of Folic Acid; 15.60 mg/L of i-Inositol; 3.02 mg/L of Niacinamide; 3.00 mg/L of Pyridoxal HCL; 0.031 mg/L of Pyridoxine HCL; 0.319 mg/L of Riboflavin; 3.17 mg/L of Thiamine HCL; 0.365 mg/L of Thymidine; and 0.680 mg/L of Vitamin B<sub>12</sub>; 25 mM of HEPES Buffer; 2.39 mg/L of Na Hypoxanthine; 15 0.105 mg/L of Lipoic Acid; 0.081 mg/L of Sodium Putrescine-2HCL; 55.0 mg/L of Sodium Pyruvate; 0.0067 mg/L of Sodium Selenite; 20uM of Ethanolamine; 0.122 mg/L of Ferric Citrate; 41.70 mg/L of Methyl-B-Cyclodextrin complexed with Linoleic Acid; 33.33 mg/L of Methyl-B-Cyclodextrin complexed with Oleic Acid; and 10 mg/L of Methyl-B-Cyclodextrin complexed with Retinal) with 2mm glutamine and 1x penstrep. (BSA (81-068-3 Bayer) 100gm dissolved in 1L DMEM for a 10% BSA stock 20 solution). Filter the media and collect 50 ul for endotoxin assay in 15ml polystyrene conical.

The transfection reaction is terminated, preferably by tag-teaming, at the end of the incubation period. Person A aspirates off the transfection media, while person B adds 1.5ml appropriate media to each well. Incubate at 37°C for 45 or 72 hours depending on the media used: 1%BSA for 45 hours or CHO-5 for 72 hours.

On day four, using a 300ul multichannel pipetter, aliquot 600ul in one 1ml deep well plate and the remaining supernatant into a 2ml deep well. The supernatants from each well can then be used in the assays described in Examples 13-20.

It is specifically understood that when activity is obtained in any of the assays described below using a supernatant, the activity originates from either the polypeptide directly (e.g., as a secreted protein) or by the polypeptide inducing expression of other proteins, which are then secreted into the supernatant. Thus, the invention further provides a method of identifying the protein in the supernatant characterized by an activity in a particular assay.

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## **Example 12: Construction of GAS Reporter Construct**

One signal transduction pathway involved in the differentiation and proliferation of cells is called the Jaks-STATs pathway. Activated proteins in the Jaks-STATs pathway bind to gamma activation site "GAS" elements or interferon-sensitive responsive element ("ISRE"), located in the promoter of many genes. The binding of a protein to these elements alter the expression of the associated gene.

GAS and ISRE elements are recognized by a class of transcription factors called Signal Transducers and Activators of Transcription, or "STATs." There are six members of the STATs family. Stat1 and Stat3 are present in many cell types, as is Stat2 (as response to IFN-alpha is widespread). Stat4 is more restricted and is not in many cell types though it has been found in T helper class I, cells after treatment with IL-12. Stat5 was originally called mammary growth factor, but has been found at higher concentrations in other cells including myeloid cells. It can be activated in tissue culture cells by many cytokines.

The STATs are activated to translocate from the cytoplasm to the nucleus upon tyrosine phosphorylation by a set of kinases known as the Janus Kinase ("Jaks") family. Jaks represent a distinct family of soluble tyrosine kinases and include Tyk2, Jak1, Jak2, and Jak3. These kinases display significant sequence similarity and are generally catalytically inactive in resting cells.

The Jaks are activated by a wide range of receptors summarized in the Table below. (Adapted from review by Schidler and Darnell, Ann. Rev. Biochem. 64:621-51 (1995).) A cytokine receptor family, capable of activating Jaks, is divided into two groups: (a) Class 1 includes receptors for IL-2, IL-3, IL-4, IL-6, IL-7, IL-9, IL-11, IL-12, IL-15, Epo, PRL, GH, G-CSF, GM-CSF, LIF, CNTF, and thrombopoietin; and (b) Class 2 includes IFN-a, IFN-g, and IL-10. The Class 1 receptors share a conserved cysteine motif (a set of four conserved cysteines and one tryptophan) and a WSXWS motif (a membrane proxial region encoding Trp-Ser-Xxx-Trp-Ser (SEQ ID NO:2)).

Thus, on binding of a ligand to a receptor, Jaks are activated, which in turn activate STATs, which then translocate and bind to GAS elements. This entire process is encompassed in the Jaks-STATs signal transduction pathway.

Therefore, activation of the Jaks-STATs pathway, reflected by the binding of the GAS or the ISRE element, can be used to indicate proteins involved in the proliferation and differentiation of cells. For example, growth factors and cytokines are known to activate the Jaks-STATs pathway. (See Table below.) Thus, by using GAS elements linked to reporter molecules, activators of the Jaks-STATs pathway can be identified.

	Ligand	tyk2	<u>JAKs</u> <u>Jak 1</u>	Jak2	Jak3	<u>STATS</u>	GAS(elements) or ISRE
5	IFN family IFN-a/B IFN-g Il-10	+	+ + ?	- + ?	- -	1,2,3 1 1,3	ISRE GAS (IRF1>Lys6>IFP)
10	gp130 family IL-6 (Pleiotrohic) Il-11(Pleiotrohic) OnM(Pleiotrohic)	+ ? ?	++++++	+ ? +	????	1,3 1,3 1,3	GAS (IRF1>Lys6>IFP)
15	LIF(Pleiotrohic) CNTF(Pleiotrohic) G-CSF(Pleiotrohic) IL-12(Pleiotrohic)	? -/+ ? +	+ + +	+ + ? +	? ? +	1,3 1,3 1,3 1,3	
20	g-C family IL-2 (lymphocytes) IL-4 (lymph/myeloid) IL-7 (lymphocytes) IL-9 (lymphocytes) IL-13 (lymphocyte) IL-15	- - - - - ?	+ + + + +	- - - ? ?	+ + + + ?	1,3,5 6 5 5 6 5	GAS GAS (IRF1 = IFP >>Ly6)(IgH) GAS GAS GAS GAS GAS
30	gp140 family IL-3 (myeloid) IL-5 (myeloid) GM-CSF (myeloid)	- -	- -	+ + +	- -	5 5 5	GAS (IRF1>IFP>>Ly6) GAS GAS
35	Growth hormone fami GH PRL EPO	? ? ?	- +/- -	+ + +	- - -	5 1,3,5 5	GAS(B-CAS>IRF1=IFP>>Ly6)
40	Receptor Tyrosine Kin EGF PDGF CSF-1	nases ? ?	+ + + +	+ + +	- - -	1,3 1,3 1,3	GAS (IRF1) GAS (not IRF1)

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To construct a synthetic GAS containing promoter element, which is used in the Biological Assays described in Examples 13-14, a PCR based strategy is employed to generate a GAS-SV40 promoter sequence. The 5' primer contains four tandem copies of the GAS binding site found in the IRF1 promoter and previously demonstrated to bind STATs upon induction with a range of cytokines (Rothman et al., Immunity 1:457-468 (1994).), although other GAS or ISRE elements can be used instead. The 5' primer also contains 18bp of sequence complementary to the SV40 early promoter sequence and is flanked with an XhoI site. The sequence of the 5' primer is: 5':GCGCCTCGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTCCCCGAAATGATTTC

The downstream primer is complementary to the SV40 promoter and is flanked with a Hind III site: 5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO:4)

PCR amplification is performed using the SV40 promoter template present in the B-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with Xhol/Hind III and subcloned into BLSK2-. (Stratagene.) Sequencing with forward and reverse primers confirms that the insert contains the following sequence:

With this GAS promoter element linked to the SV40 promoter, a GAS:SEAP2 reporter construct is next engineered. Here, the reporter molecule is a secreted alkaline phosphatase, or "SEAP." Clearly, however, any reporter molecule can be instead of SEAP, in this or in any of the other Examples. Well known reporter molecules that can be used instead of SEAP include chloramphenicol acetyltransferase (CAT), luciferase, alkaline phosphatase, B-galactosidase, green fluorescent protein (GFP), or any protein detectable by an antibody.

The above sequence confirmed synthetic GAS-SV40 promoter element is subcloned into the pSEAP-Promoter vector obtained from Clontech using HindIII and XhoI, effectively replacing the SV40 promoter with the amplified GAS:SV40 promoter element, to create the GAS-SEAP vector. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

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Thus, in order to generate mammalian stable cell lines expressing the GAS-SEAP reporter, the GAS-SEAP cassette is removed from the GAS-SEAP vector using SalI and NotI, and inserted into a backbone vector containing the neomycin resistance gene, such as pGFP-1 (Clontech), using these restriction sites in the multiple cloning site, to create the GAS-SEAP/Neo vector. Once this vector is transfected into mammalian cells, this vector can then be used as a reporter molecule for GAS binding as described in Examples 13-14.

Other constructs can be made using the above description and replacing GAS with a different promoter sequence. For example, construction of reporter molecules containing NFK-B and EGR promoter sequences are described in Examples 15 and 16. However, many other promoters can be substituted using the protocols described in these Examples. For instance, SRE, IL-2, NFAT, or Osteocalcin promoters can be substituted, alone or in combination (e.g., GAS/NF-KB/EGR, GAS/NF-KB, Il-2/NFAT, or NF-KB/GAS). Similarly, other cell lines can be used to test reporter construct activity, such as HELA (epithelial), HUVEC (endothelial), Reh (B-cell), Saos-2 (osteoblast), HUVAC (aortic), or Cardiomyocyte.

## Example 13: High-Throughput Screening Assay for T-cell Activity.

The following protocol is used to assess T-cell activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate T-cells. T-cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The T-cell used in this assay is Jurkat T-cells (ATCC Accession No. TIB-152), although Molt-3 cells (ATCC Accession No. CRL-1552) and Molt-4 cells (ATCC Accession No. CRL-1582) cells can also be used.

Jurkat T-cells are lymphoblastic CD4+ Th1 helper cells. In order to generate stable cell lines, approximately 2 million Jurkat cells are transfected with the GAS-SEAP/neo vector using DMRIE-C (Life Technologies)(transfection procedure described below). The transfected cells are seeded to a density of approximately 20,000 cells per well and transfectants resistant to 1 mg/ml genticin selected. Resistant colonies are expanded and then tested for their response to increasing concentrations of interferon gamma. The dose response of a selected clone is demonstrated.

Specifically, the following protocol will yield sufficient cells for 75 wells containing 200 ul of cells. Thus, it is either scaled up, or performed in multiple to generate sufficient cells for multiple 96 well plates. Jurkat cells are maintained in RPMI + 10% serum with 1%Pen-Strep. Combine 2.5 mls of OPTI-MEM (Life Technologies)

WO 98/54963

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with 10 ug of plasmid DNA in a T25 flask. Add 2.5 ml OPTI-MEM containing 50 ul of DMRIE-C and incubate at room temperature for 15-45 mins.

During the incubation period, count cell concentration, spin down the required number of cells ( $10^7$  per transfection), and resuspend in OPTI-MEM to a final concentration of  $10^7$  cells/ml. Then add 1ml of 1 x  $10^7$  cells in OPTI-MEM to T25 flask and incubate at 37°C for 6 hrs. After the incubation, add 10 ml of RPMI + 15% serum.

The Jurkat:GAS-SEAP stable reporter lines are maintained in RPMI + 10% serum, 1 mg/ml Genticin, and 1% Pen-Strep. These cells are treated with supernatants containing a polypeptide as produced by the protocol described in Example 11.

On the day of treatment with the supernatant, the cells should be washed and resuspended in fresh RPMI + 10% serum to a density of 500,000 cells per ml. The exact number of cells required will depend on the number of supernatants being screened. For one 96 well plate, approximately 10 million cells (for 10 plates, 100 million cells) are required.

Transfer the cells to a triangular reservoir boat, in order to dispense the cells into a 96 well dish, using a 12 channel pipette. Using a 12 channel pipette, transfer 200 ul of cells into each well (therefore adding 100, 000 cells per well).

After all the plates have been seeded, 50 ul of the supernatants are transferred directly from the 96 well plate containing the supernatants into each well using a 12 channel pipette. In addition, a dose of exogenous interferon gamma (0.1, 1.0, 10 ng) is added to wells H9, H10, and H11 to serve as additional positive controls for the assay.

The 96 well dishes containing Jurkat cells treated with supernatants are placed in an incubator for 48 hrs (note: this time is variable between 48-72 hrs). 35 ul samples from each well are then transferred to an opaque 96 well plate using a 12 channel pipette. The opaque plates should be covered (using sellophene covers) and stored at -20°C until SEAP assays are performed according to Example 17. The plates containing the remaining treated cells are placed at 4°C and serve as a source of material for repeating the assay on a specific well if desired.

As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate Jurkat T cells. Over 30 fold induction is typically observed in the positive control wells.

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## Example 14: High-Throughput Screening Assay Identifying Myeloid Activity

The following protocol is used to assess myeloid activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate myeloid cells. Myeloid cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The myeloid cell used in this assay is U937, a pre-monocyte cell line, although TF-1, HL60, or KG1 can be used.

To transiently transfect U937 cells with the GAS/SEAP/Neo construct produced in Example 12, a DEAE-Dextran method (Kharbanda et. al., 1994, Cell Growth & Differentiation, 5:259-265) is used. First, harvest 2x10e<sup>7</sup> U937 cells and wash with PBS. The U937 cells are usually grown in RPMI 1640 medium containing 10% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 mg/ml streptomycin.

Next, suspend the cells in 1 ml of 20 mM Tris-HCl (pH 7.4) buffer containing 0.5 mg/ml DEAE-Dextran, 8 ug GAS-SEAP2 plasmid DNA, 140 mM NaCl, 5 mM KCl, 375 uM Na<sub>2</sub>HPO<sub>4</sub>.7H<sub>2</sub>O, 1 mM MgCl<sub>2</sub>, and 675 uM CaCl<sub>2</sub>. Incubate at 37°C for 45 min.

Wash the cells with RPMI 1640 medium containing 10% FBS and then resuspend in 10 ml complete medium and incubate at 37°C for 36 hr.

The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 400 ug/ml G418 for couple of passages.

These cells are tested by harvesting  $1x10^8$  cells (this is enough for ten 96-well plates assay) and wash with PBS. Suspend the cells in 200 ml above described growth medium, with a final density of  $5x10^5$  cells/ml. Plate 200 ul cells per well in the 96-well plate (or  $1x10^5$  cells/well).

Add 50 ul of the supernatant prepared by the protocol described in Example 11. Incubate at 37°C for 48 to 72 hr. As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate U937 cells. Over 30 fold induction is typically observed in the positive control wells. SEAP assay the supernatant according to the protocol described in Example 17.

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## Example 15: High-Throughput Screening Assay Identifying Neuronal Activity.

When cells undergo differentiation and proliferation, a group of genes are activated through many different signal transduction pathways. One of these genes, EGR1 (early growth response gene 1), is induced in various tissues and cell types upon activation. The promoter of EGR1 is responsible for such induction. Using the EGR1 promoter linked to reporter molecules, activation of cells can be assessed.

Particularly, the following protocol is used to assess neuronal activity in PC12 cell lines. PC12 cells (rat phenochromocytoma cells) are known to proliferate and/or differentiate by activation with a number of mitogens, such as TPA (tetradecanoyl phorbol acetate), NGF (nerve growth factor), and EGF (epidermal growth factor). The EGR1 gene expression is activated during this treatment. Thus, by stably transfecting PC12 cells with a construct containing an EGR promoter linked to SEAP reporter, activation of PC12 cells can be assessed.

The EGR/SEAP reporter construct can be assembled by the following protocol. The EGR-1 promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene 6:867-871 (1991)) can be PCR amplified from human genomic DNA using the following primers:

- 5' GCGCTCGAGGGATGACAGCGATAGAACCCCGG -3' (SEQ ID NO:6)
- 5' GCGAAGCTTCGCGACTCCCCGGATCCGCCTC-3' (SEQ ID NO:7)

Using the GAS:SEAP/Neo vector produced in Example 12, EGR1 amplified product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector using restriction enzymes XhoI/HindIII, removing the GAS/SV40 stuffer. Restrict the EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1 promoter.

To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30 dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and allowed to air dry for 2 hr.

PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker) containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is done every three to four days. Cells are removed from the plates by scraping and resuspended with pipetting up and down for more than 15 times.

Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine protocol described in Example 11. EGR-SEAP/PC12 stable cells are obtained by growing the cells in 300 ug/ml G418. The G418-free medium is used for routine

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growth but every one to two months, the cells should be re-grown in 300 ug/ml G418 for couple of passages.

To assay for neuronal activity, a 10 cm plate with cells around 70 to 80% confluent is screened by removing the old medium. Wash the cells once with PBS (Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as  $5x10^5$  cells/ml.

Add 200 ul of the cell suspension to each well of 96-well plate (equivalent to  $1 \times 10^5$  cells/well). Add 50 ul supernatant produced by Example 11, 37°C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ul of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 17.

## Example 16: High-Throughput Screening Assay for T-cell Activity

NF-κB (Nuclear Factor κB) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by expression of certain viral gene products. As a transcription factor, NF-κB regulates the expression of genes involved in immune cell activation, control of apoptosis (NF-κB appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

In non-stimulated conditions, NF-  $\kappa B$  is retained in the cytoplasm with I- $\kappa B$  (Inhibitor  $\kappa B$ ). However, upon stimulation, I-  $\kappa B$  is phosphorylated and degraded, causing NF-  $\kappa B$  to shuttle to the nucleus, thereby activating transcription of target genes. Target genes activated by NF-  $\kappa B$  include IL-2, IL-6, GM-CSF, ICAM-1 and class 1 MHC.

Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF-kB promoter element are used to screen the supernatants produced in Example 11. Activators or inhibitors of NF-kB would be useful in treating

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diseases. For example, inhibitors of NF-kB could be used to treat those diseases related to the acute or chronic activation of NF-kB, such as rheumatoid arthritis.

To construct a vector containing the NF-κB promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF-κB binding site (GGGGACTTTCCC) (SEQ ID NO:8), 18 bp of sequence complementary to the 5' end of the SV40 early promoter sequence, and is flanked with an XhoI site: 5':GCGGCCTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCATCTCAATTAG:3' (SEQ ID NO:9)

The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:

5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO:4)

PCR amplification is performed using the SV40 promoter template present in the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2-. (Stratagene)

Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

5':CTCGAGGGACTTTCCCGGGGACTTTCCGGGACTTTCC
ATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCCCTAACTCCGCCCA
TCCCGCCCCTAACTCCGCCCAGTTCCGCCCATTCTCCGCCCCATGGCTGACT
AATTTTTTTATITATGCAGAGGCCGAGGCCGCCTCGGCCTCTGAGCTATTC
CAGAAGTAGTGAGGAGGCCTTTTTTTGGAGGCCTAGGCTTTTGCAAAAAGCTT:
3' (SEQ ID NO:10)

Next, replace the SV40 minimal promoter element present in the pSEAP2promoter plasmid (Clontech) with this NF-kB/SV40 fragment using XhoI and HindIII.
However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

In order to generate stable mammalian cell lines, the NF-κB/SV40/SEAP

cassette is removed from the above NF-κB/SEAP vector using restriction enzymes SalI and NotI, and inserted into a vector containing neomycin resistance. Particularly, the NF-κB/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP gene, after restricting pGFP-1 with SalI and NotI.

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Once NF-kB/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are created and maintained according to the protocol described in Example 13. Similarly, the method for assaying supernatants with these stable Jurkat T-cells is also described in Example 13. As a positive control, exogenous TNF alpha (0.1,1, 10 ng) is added to wells H9, H10, and H11, with a 5-10 fold activation typically observed.

### **Example 17: Assay for SEAP Activity**

As a reporter molecule for the assays described in Examples 13-16, SEAP activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the following general procedure. The Tropix Phospho-light Kit supplies the Dilution, Assay, and Reaction Buffers used below.

Prime a dispenser with the 2.5x Dilution Buffer and dispense 15  $\mu$ l of 2.5x dilution buffer into Optiplates containing 35  $\mu$ l of a supernatant. Seal the plates with a plastic sealer and incubate at 65°C for 30 min. Separate the Optiplates to avoid uneven heating.

Cool the samples to room temperature for 15 minutes. Empty the dispenser and prime with the Assay Buffer. Add 50 µl Assay Buffer and incubate at room temperature 5 min. Empty the dispenser and prime with the Reaction Buffer (see the table below). Add 50 µl Reaction Buffer and incubate at room temperature for 20 minutes. Since the intensity of the chemiluminescent signal is time dependent, and it takes about 10 minutes to read 5 plates on luminometer, one should treat 5 plates at each time and start the second set 10 minutes later.

Read the relative light unit in the luminometer. Set H12 as blank, and print the results. An increase in chemiluminescence indicates reporter activity.

Reaction Buffer Formulation:

Acception D	unci l'ormulation.	
# of plates	Rxn buffer diluent (ml)	CSPD (ml)
10	60	3
11	65	3.25
12	70 ·	3.5
13	75	3.75
14	80	4
15	85	4.25
16	90	4.5
<b>17</b>	95	4.75
18	100	5
19	105	5.25
20	110	5.5
21	115	5.75
22	120	6

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23	125	6.25
24	130	6.5
25	135	6.75
26	140	7
27	145	7.25
28	150	7.5
29	155	7.75
30	160	8
31	165	8.25
32	170	8.5
33	175	8.75
34	180	9
35	185	9.25
36	190	9.5
37	195	9.75
38	200	10
39	205	10.25
40	210	10.5
41	215	10.75
42	<b>220</b> .	11
43	225	11.25
44	230	11.5
45	235	11.75
46	240	12
47	245	12.25
48	250	12.5
49	255	12.75
50	260	13

# Example 18: High-Throughput Screening Assay Identifying Changes in Small Molecule Concentration and Membrane Permeability

Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane potential. These alterations can be measured in an assay to identify supernatants which bind to receptors of a particular cell. Although the following protocol describes an assay for calcium, this protocol can easily be modified to detect changes in potassium, sodium, pH, membrane potential, or any other small molecule which is detectable by a fluorescent probe.

The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small molecules. Clearly, any fluorescent molecule detecting a small molecule can be used instead of the calcium fluorescent molecule, fluo-3, used here.

For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black 96-well plate with clear bottom. The plate is incubated in a CO<sub>2</sub> incubator for 20 hours. The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash.

A stock solution of 1 mg/ml fluo-3 is made in 10% pluronic acid DMSO. To load the cells with fluo-3, 50 ul of 12 ug/ml fluo-3 is added to each well. The plate is incubated at  $37^{\circ}$ C in a  $CO_2$  incubator for 60 min. The plate is washed four times in the Biotek washer with HBSS leaving 100 ul of buffer.

For non-adherent cells, the cells are spun down from culture media. Cells are re-suspended to 2-5x10<sup>6</sup> cells/ml with HBSS in a 50-ml conical tube. 4 ul of 1 mg/ml fluo-3 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. The tube is then placed in a 37°C water bath for 30-60 min. The cells are washed twice with HBSS, resuspended to 1x10<sup>6</sup> cells/ml, and dispensed into a microplate, 100 ul/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley CellWash with 200 ul, followed by an aspiration step to 100 ul final volume.

For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-3. The supernatant is added to the well, and a change in fluorescence is detected.

To measure the fluorescence of intracellular calcium, the FLIPR is set for the following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and (6) Sample addition is 50 ul. Increased emission at 530 nm indicates an extracellular signaling event which has resulted in an increase in the intracellular Ca<sup>++</sup> concentration.

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## Example 19: High-Throughput Screening Assay Identifying Tyrosine Kinase Activity

The Protein Tyrosine Kinases (PTK) represent a diverse group of transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase RPTK) group are receptors for a range of mitogenic and metabolic growth factors including the PDGF, FGF, EGF, NGF, HGF and Insulin receptor subfamilies. In addition there are a large family of RPTKs for which the corresponding ligand is unknown. Ligands for RPTKs include mainly secreted small proteins, but also membrane-bound and extracellular matrix proteins.

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Activation of RPTK by ligands involves ligand-mediated receptor dimerization, resulting in transphosphorylation of the receptor subunits and activation of the cytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and non-receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members of which mediate signal transduction triggered by the cytokine superfamily of receptors (e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

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Because of the wide range of known factors capable of stimulating tyrosine kinase activity, the identification of novel human secreted proteins capable of activating tyrosine kinase signal transduction pathways are of interest. Therefore, the following protocol is designed to identify those novel human secreted proteins capable of activating the tyrosine kinase signal transduction pathways.

Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyne Silent Screen Plates purchased from Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 100% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or 10% Matrigel purchased from Becton Dickinson (Bedford,MA), or calf serum, rinsed with PBS and stored at 4°C. Cell growth on these plates is assayed by seeding 5,000 cells/well in growth medium and indirect quantitation of cell number through use of alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford,MA) are used to cover the Loprodyne Silent Screen Plates. Falcon Microtest III cell culture plates can also be used in some proliferation experiments.

To prepare extracts, A431 cells are seeded onto the nylon membranes of Loprodyne plates (20,000/200ml/well) and cultured overnight in complete medium. Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example 11, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na3VO4, 2 mM Na4P2O7 and a cocktail of protease inhibitors (# 1836170) obtained from Boeheringer Mannheim (Indianapolis, IN) is added to each well and the plate is shaken on a rotating shaker for 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract filtered through the 0.45 mm membrane bottoms of each well using house vacuum. Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum manifold and immediately placed on ice. To obtain extracts clarified by centrifugation, the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4°C at 16,000 x g.

Test the filtered extracts for levels of tyrosine kinase activity. Although many methods of detecting tyrosine kinase activity are known, one method is described here.

Generally, the tyrosine kinase activity of a supernatant is evaluated by determining its ability to phosphorylate a tyrosine residue on a specific substrate (a

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biotinylated peptide). Biotinylated peptides that can be used for this purpose include PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and PSK2 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for a range of tyrosine kinases and are available from Boehringer Mannheim.

The tyrosine kinase reaction is set up by adding the following components in order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg<sub>2+</sub> (5mM ATP/50mM MgCl<sub>2</sub>), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride, pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl<sub>2</sub>, 5 mM MnCl<sub>2</sub>, 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the components gently and preincubate the reaction mix at 30°C for 2 min. Initial the reaction by adding 10ul of the control enzyme or the filtered supernatant.

The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mm EDTA and place the reactions on ice.

Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction mixture to a microtiter plate (MTP) module and incubating at 37°C for 20 min. This allows the streptavadin coated 96 well plate to associate with the biotinylated peptide. Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of anti-phospotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-POD(0.5u/ml)) to each well and incubate at 37°C for one hour. Wash the well as above.

Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and incubate at room temperature for at least 5 mins (up to 30 min). Measure the absorbance of the sample at 405 nm by using ELISA reader. The level of bound peroxidase activity is quantitated using an ELISA reader and reflects the level of tyrosine kinase activity.

# Example 20: High-Throughput Screening Assay Identifying Phosphorylation Activity

As a potential alternative and/or compliment to the assay of protein tyrosine
kinase activity described in Example 19, an assay which detects activation
(phosphorylation) of major intracellular signal transduction intermediates can also be
used. For example, as described below one particular assay can detect tyrosine
phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other
molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase,
Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other

phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by substituting these molecules for Erk-1 or Erk-2 in the following assay.

Specifically, assay plates are made by coating the wells of a 96-well ELISA plate with 0.1ml of protein G (lug/ml) for 2 hr at room temp, (RT). The plates are then rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates are then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1 and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this step can easily be modified by substituting a monoclonal antibody detecting any of the above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4°C until use.

A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and cultured overnight in growth medium. The cells are then starved for 48 hr in basal medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants obtained in Example 11 for 5-20 minutes. The cells are then solubilized and extracts filtered directly into the assay plate.

After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a positive control, a commercial preparation of MAP kinase (10ng/well) is used in place of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody (1ug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The bound polyclonal antibody is then quantitated by successive incubations with Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over background indicates a phosphorylation.

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## Example 21: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is be isolated. cDNA is then generated from these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in SEQ ID NO:X. Suggested PCR conditions consist of 35 cycles at 95°C for 30 seconds; 60-120 seconds at 52-58°C; and 60-120 seconds at 70°C, using buffer solutions described in Sidransky, D., et al., Science 252:706 (1991).

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PCR products are then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase. (Epicentre Technologies).

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The intron-exon borders of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations is then cloned and sequenced to validate the results of the direct sequencing.

PCR products is cloned into T-tailed vectors as described in Holton, T.A. and Graham, M.W., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals are identified by mutations not present in unaffected individuals.

Genomic rearrangements are also observed as a method of determining alterations in a gene corresponding to a polynucleotide. Genomic clones isolated according to Example 2 are nick-translated with digoxigenindeoxy-uridine 5'triphosphate (Boehringer Manheim), and FISH performed as described in Johnson, Cg. et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the corresponding genomic locus.

15 Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson, Cv. 20 et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosome alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and translocations. These alterations are used as a diagnostic marker for an associated disease.

## Example 22: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

For example, antibody-sandwich ELISAs are used to detect polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method described in Example 10.

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The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, serial dilutions of the sample should be used to validate results. The plates are then washed three times with deionized or distilled water to remove unbounded polypeptide.

Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove unbounded conjugate.

Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale). Interpolate the concentration of the polypeptide in the sample using the standard curve.

#### Example 23: Formulating a Polypeptide

The secreted polypeptide composition will be formulated and dosed in a fashion consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with the secreted polypeptide alone), the site of delivery, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

As a general proposition, the total pharmaceutically effective amount of secreted polypeptide administered parenterally per dose will be in the range of about 1  $\mu$ g/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the secreted polypeptide is typically administered at a dose rate of about 1  $\mu$ g/kg/hour to about 50  $\mu$ g/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending on the desired effect.

Pharmaceutical compositions containing the secreted protein of the invention are administered orally, rectally, parenterally, intracistemally, intravaginally,

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intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), bucally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

The secreted polypeptide is also suitably administered by sustained-release systems. Suitable examples of sustained-release compositions include semi-permeable polymer matrices in the form of shaped articles, e.g., films, or mirocapsules. Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman, U. et al., Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (R. Langer et al., J. Biomed. Mater. Res. 15:167-277 (1981), and R. Langer, Chem. Tech. 12:98-105 (1982)), ethylene vinyl acetate (R. Langer et al.) or poly-D- (-)-3-hydroxybutyric acid (EP 133,988). Sustained-release compositions also include liposomally entrapped polypeptides. Liposomes containing the secreted polypeptide are prepared by methods known per se: DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. USA 82:3688-3692 (1985); Hwang et al., Proc. Natl. Acad. Sci. USA 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal secreted polypeptide therapy.

For parenteral administration, in one embodiment, the secreted polypeptide is formulated generally by mixing it at the desired degree of purity, in a unit dosage injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations employed and is compatible with other ingredients of the formulation. For example, the formulation preferably does not include oxidizing agents and other compounds that are known to be deleterious to polypeptides.

Generally, the formulations are prepared by contacting the polypeptide uniformly and intimately with liquid carriers or finely divided solid carriers or both. Then, if necessary, the product is shaped into the desired formulation. Preferably the carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood of the recipient. Examples of such carrier vehicles include water, saline, Ringer's solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl oleate are also useful herein, as well as liposomes.

WO 98/54963 PCT/US98/11422

259

The carrier suitably contains minor amounts of additives such as substances that enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at the dosages and concentrations employed, and include buffers such as phosphate, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, manose, or dextrins; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates, poloxamers, or PEG.

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The secreted polypeptide is typically formulated in such vehicles at a concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.

Any polypeptide to be used for therapeutic administration can be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutic polypeptide compositions generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

Polypeptides ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous polypeptide solution, and the resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the lyophilized polypeptide using bacteriostatic Water-for-Injection.

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the present invention may be employed in conjunction with other therapeutic compounds.

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## Example 24: Method of Treating Decreased Levels of the Polypeptide

It will be appreciated that conditions caused by a decrease in the standard or normal expression level of a secreted protein in an individual can be treated by administering the polypeptide of the present invention, preferably in the secreted form. Thus, the invention also provides a method of treatment of an individual in need of an increased level of the polypeptide comprising administering to such an individual a pharmaceutical composition comprising an amount of the polypeptide to increase the activity level of the polypeptide in such an individual.

For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the polypeptide for six consecutive days. Preferably, the polypeptide is in the secreted form. The exact details of the dosing scheme, based on administration and formulation, are provided in Example 23.

## Example 25: Method of Treating Increased Levels of the Polypeptide

Antisense technology is used to inhibit production of a polypeptide of the present invention. This technology is one example of a method of decreasing levels of a polypeptide, preferably a secreted form, due to a variety of etiologies, such as cancer.

For example, a patient diagnosed with abnormally increased levels of a polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, 2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is provided in Example 23.

### Example 26: Method of Treatment Using Gene Therapy

One method of gene therapy transplants fibroblasts, which are capable of expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and separated into small pieces. Small chunks of the tissue are placed on a wet surface of a tissue culture flask, approximately ten pieces are placed in each flask. The flask is turned upside down, closed tight and left at room temperature over night. After 24 hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to the bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS, penicillin and streptomycin) is added. The flasks are then incubated at 37°C for approximately one week.

At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long terminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is fractionated on agarose gel and purified, using glass beads.

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The cDNA encoding a polypeptide of the present invention can be amplified using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to transform bacteria HB101, which are then plated onto agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media, containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is produced.

The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

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## Example 27: Method of Treatment Using Gene Therapy - In Vivo

Another aspect of the present invention is using *in vivo* gene therapy methods to treat disorders, diseases and conditions. The gene therapy method relates to the introduction of naked nucleic acid (DNA, RNA, and antisense DNA or RNA) sequences into an animal to increase or decrease the expression of the polypeptide of the present invention. A polynucleotide of the present invention may be operatively linked to a promoter or any other genetic elements necessary for the expression of the encoded polypeptide by the target tissue. Such gene therapy and delivery techniques and methods are known in the art, see, for example, WO90/11092, WO98/11779; U.S. Patent NO. 5693622, 5705151, 5580859; Tabata H. et al. (1997) Cardiovasc. Res. 35(3):470-479, Chao J et al. (1997) Pharmacol. Res. 35(6):517-522, Wolff J.A. (1997) Neuromuscul. Disord. 7(5):314-318, Schwartz B. et al. (1996) Gene Ther. 3(5):405-411, Tsurumi Y. et al. (1996) Circulation 94(12):3281-3290 (incorporated herein by reference).

The polynucleotide constructs of the present invention may be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, intestine and the like). These polynucleotide constructs can be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

The term "naked" polynucleotide, DNA or RNA, refers to sequences that are free from any delivery vehicle that acts to assist, promote, or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotides may also be delivered in liposome formulations (such as those taught in Felgner P.L. et al. (1995) Ann. NY Acad. Sci. 772:126-139 and Abdallah B. et al. (1995) Biol. Cell 85(1):1-7) which can be prepared by methods well known to those skilled in the art.

The polynucleotide vector constructs of the present invention used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Any strong promoter known to those skilled in the art can be used for driving the expression of DNA. Unlike other gene therapies techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.

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The polynucleotide construct of the present invention can be delivered to the interstitial space of tissues within the an animal, including of muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and connective tissue. Interstitial space of the tissues comprises the intercellular fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less 15 completely differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. In vivo muscle cells are particularly competent in their ability to take up and express polynucleotides.

For the naked polynucleotide injection, an effective dosage amount of DNA or RNA will be in the range of from about 0.05 g/kg body weight to about 50 mg/kg body weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration. The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial tissues, throat or mucous membranes of the nose. In addition, naked polynucleotide constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

The dose response effects of injected polynucleotide in muscle in vivo is determined as follows. Suitable template DNA for production of mRNA coding for the polypeptide of the present invention is prepared in accordance with a standard recombinant DNA methodology. The template DNA, which may be either circular or linear, is either used as naked DNA or complexed with

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liposomes. The quadriceps muscles of mice are then injected with various amounts of the template DNA.

Five to six week old female and male Balb/C mice are anesthetized by intraperitoneal injection with 0.3 ml of 2.5% Avertin. A 1.5 cm incision is made on the anterior thigh, and the quadriceps muscle is directly visualized. The template DNA is injected in 0.1 ml of carrier in a 1 cc syringe through a 27 gauge needle over one minute, approximately 0.5 cm from the distal insertion site of the muscle into the knee and about 0.2 cm deep. A suture is placed over the injection site for future localization, and the skin is closed with stainless steel clips.

After an appropriate incubation time (e.g., 7 days) muscle extracts are prepared by excising the entire quadriceps. Every fifth 15 um cross-section of the individual quadriceps muscles is histochemically stained for protein expression. A time course for protein expression may be done in a similar fashion except that quadriceps from different mice are harvested at different times. Persistence of DNA in muscle following injection may be determined by Southern blot analysis after preparing total cellular DNA and HIRT supernatants from injected and control mice. The results of the above experimentation in mice can be use to extrapolate proper dosages and other treatment parameters in humans and other animals using naked DNA of the present invention.

It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

The entire disclosure of each document cited (including patents, patent applications, journal articles, abstracts, laboratory manuals, books, or other disclosures) in the Background of the Invention, Detailed Description, and Examples is hereby incorporated herein by reference.

## Sequence Listing

(1) GENERAL INFORMATION: (i) APPLICANT: Human Genome Sciences, Inc., et al. 5 (ii) TITLE OF INVENTION: 207 Human Secreted Proteins 10 (iii) NUMBER OF SEQUENCES: 800 (iv) CORRESPONDENCE ADDRESS: 15 (A) ADDRESSEE: Human Genome Sciences, Inc. (B) STREET: 9410 Key West Avenue 20 (C) CITY: Rockville (D) STATE: Maryland (E) COUNTRY: USA 25 (F) ZIP: 20850 30 (v) COMPUTER READABLE FORM: (A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage (B) COMPUTER: HP Vectra 486/33 35 (C) OPERATING SYSTEM: MSDOS version 6.2 (D) SOFTWARE: ASCII Text 40 (vi) CURRENT APPLICATION DATA: (A) APPLICATION NUMBER: 45 (B) FILING DATE: (C) CLASSIFICATION: 50 (vii) PRIOR APPLICATION DATA: (A) APPLICATION NUMBER: 55 (B) FILING DATE:

	(VIII) Alloway, mark an obtained in	
5	(A) NAME: Kenley K. Hoover	
	(B) REGISTRATION NUMBER: 40,302	
	(C) REFERENCE/DOCKET NUMBER: PZ007PCT	·
10		
	(vi) TELECOMMUNICATION INFORMATION:	
1.5	(A) TELEPHONE: (301) 309-8504	
15	(B) TELEFAX: (301) 309-8439	
20	TO NO. 1.	
	(2) INFORMATION FOR SEQ ID NO: 1:	
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 733 base pairs	
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
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	TCAAGTTCAA CTGGTACGTG GACGGCGTGG AGGTGCATAA TGCCAAGACA AAGCCGCGGG	240
40	AGGAGCAGTA CAACAGCACG TACCGTGTGG TCAGCGTCCT CACCGTCCTG CACCAGGACT	300
	GGCTGAATGG CAAGGAGTAC AAGTGCAAGG TCTCCAACAA AGCCCTCCCA ACCCCCATCG	360
	AGAAAACCAT CTCCAAAGCC AAAGGGCAGC CCCGAGAACC ACAGGTGTAC ACCCTGCCCC	420
45	CATCCCGGGA TGAGCTGACC AAGAACCAGG TCAGCCTGAC CTGCCTGGTC AAAGGCTTCT	480
	ATCCAAGCGA CATCGCCGTG GAGTGGGAGA GCAATGGGCA GCCGGAGAAC AACTACAAGA	540
50	CCACGCCTCC CGTGCTGGAC TCCGACGGCT CCTTCTTCCT CTACAGCAAG CTCACCGTGG	600
	ACAAGAGCAG GTGGCAGCAG GGGAACGTCT TCTCATGCTC CGTGATGCAT GAGGCTCTGC	660
	ACAACCACTA CACGCAGAAG AGCCTCTCCC TGTCTCCGGG TAAATGAGTG CGACGGCCGC	720
55	GACTCTAGAG GAT	733

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	(B) TYPE: amino acid	
5	(D) TOPOLOGY: linear	
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	(	
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10	· ·	
10	1 5	
	·	
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	(B) TYPE: nucleic acid	
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	(D) TOPOLOGY: linear	
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50		
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	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:	
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45		
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	(4) = = = = = = = = = = = = = = = = = = =	
•	(i) SEQUENCE CHARACTERISTICS:	
50	(A) LENGTH: 271 base pairs	
50	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
	(D) TOPOLOGI: Timean	
E	A CONTROL DESCRIPTION OF YOUR E.	
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	AAATATCTGC CATCTCAATT AGTCAGCAAC CATAGTCCCG CCCCTAACTC CGCCCATCCC	120
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	GCCCCTAACT CCGCCCAGTT CCGCCCCATTC TCCGCCCCAT GGCTGACTAA TTTTTTTTAT	180
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5	TTTTGGAGGC CTAGGCTTTT GCAAAAAGCT T	271
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	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:	
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25	(2) INFORMATION FOR SEQ ID NO: 7:	
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 31 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:	
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	(i) SEQUENCE CHARACTERISTICS:	
45	<ul><li>(A) LENGTH: 12 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li><li>(D) TOPOLOGY: linear</li></ul>	
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	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
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	(D) TOPOLOGY: linear	
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	AGTTTTTAAG CAAGGCATGG GGAACAGGGA ATAGAACCTT TCAAAGAGGT TGCCCAGAGA	36
	AAAGCTGGGC CTCTTGCATT CGGCTTCCTT GGAGCAGCCT CTTCTGGCAG AAAGCCATCA	·42
۲0	CONCORDAD CARGOTTO DECECTA CON TOTAL CARGOTTA CTTACTACTACTACTACTACTACTACTACTACTACTACTA	48

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5	CCTGAGATGA TTCAGAACAA ATCATGCTAA CTTTGAATCC ATCCAGCCAC TTGCAAATGA	600
	TAATCAGAAG TCAGCTTGTT CACTGTTAGA AAGAAACTAA CAAAAGAGAA CCCAGAGCAA	660
	TCTAGAATCT TTGAGTGCTT GGCTTTCCAA GGATACTGCG GAGACTCTGG CCAAGCTGAT	720
10	GAMCTICIGA ARIGICACIG GCACCATATG CAACAAGAAC CACCATICAC IGAGIAGCIA	780
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	AGAGGACCGG TTGCAGCTTA CCCAGAACCA CTCCTCCAGG AGAGCTGGAT GTTTTGCGTG	900
15	CAACACCTTG AGCACTGACT GCTATTGTTC AAAAAAAGCC TTTGCTGCAT TCGGAGGACT	960
	GCCCCGTGCC CTGAGGTGAC TTCCTAACTA TGTGGTTTCA TTAGCGAATT TATTTTTTGT	1020
20	GCTGGGTGGA CATTTGTATT TTGTTAGGTT GCTGTTTTAAG CTCAAGTTTG CTGTGCTCTC	1080
	TGCAGCTACA AAACATCTTG GCATATTTAA GAKTGGCTTT TATAAATAGC TTTATTCTGA	1140
05	TATTAATCAG ATTCCCAACT TTACTGAGAA TTAAGGACTG GGGTACTTTA AAGAAATGCA	1200
25	AATAGCAATT GAAGAACCAC TECTECAGGT GGTAGCCCTG GCTAGACTGA ATTACACTAG	1260
	AAATCAGCCA GAAGGAAGCG TCCTTGGGAT CCCAGATCAC TCTTTTTTTT TTTTTTTTA	1320
30	AAAGGGGCAG CCCCTTGATG GCTCATCTCT CTGAATAACA GTTACGTCTT CATATCGATA	1380
٠	CCAGATGCCT TCTTCATCAT GCCACTGAAG CCACTCACCA CCTTCAAGAA CATGCCAACC	1440
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35	AGGTCCAAGT GGACTCTACA GAGTGCTTGA CCTCAACACA CTGGATTCCA GGTGGACTGG	1560
	ACCAAGAGCA GGCAAAGACA CGGGAACTGA AAAACTCCAC AGGGTTTGGA GAATAGAAAT	1620
40	GAAAAGCCAC GTCATATAAC TCAAGAATAA ATGGTGTTTT GGAAATTTTA AAATTATCAT	1680
	CGAAGGTGGT GAAACTATTT CAGGCCCAAA TGAAAGGAAA TCGCCAGTTG GGGATGAAAT	1740
	CACAGAGCCT GTGTTTTATG ATATGGTTGG ATGTCCACTG ATGAAATTTT AAAGGAGTTT	1800
45	CATTTTAAA AGIGCGCATG ATTCTACATA TGAGAATTCT TTAGGCCAAG AAACTGTCCT	1860
	TGGCTCAGAG GTGTTGGGAA TTAAAGCAGA GAGAAGCCAT TCGTGATGCT TAGAACCAAG	1920
50	GATGGTCATG TACACAAAGA CCATCGAGAC GGCCATTCTT GTTTACAAAA CACTTACCAA	1980
	GAAAGCACTT TGTAGGGGAA CTTTAGTAAG TTCTTCTCAT TTCATTATGT TTCTTCCAAG	2040
	GAAACAGGAG AGACTGAATT AATAATTCTC TCTTTCCTCT TAAGCACTTT TAAAATAATA	2100
55	AAGTACATCT TGAAATTTGG GGGGGCATCT CTGATTTAAA AAAAGAAAAA GGCTGCTTGA	2160
	TGTATGTTAT GCAGAGACAC TCTGCCTCTG GTGGCTGCAG AGCAATACCC AAGCCTCATT	2220
60	TOGAAGGCTC AACATTTGGA ATTGCACTTT AATTGATTAA TCCTCAATTC ATGTGGCCTT	228

0	TACCCA		. '				2526
5	TAAATAAAGT	AAACACTATG	ACATTTAAAA	АААААААА	AAAACTCGAG	GGGGGCCCGG .	2520
	ATATTATTTT	ATAGTGTCTG	CCATGCCATG	TGGAAATACT	TTATTTTTAA	CCTCAGGATT	2460
	AÇATCAAATA	CCAGCACCCC	ACCTGCACAA	TGGGGGTGGA	AAACTTTTGT	ATCCCTAAGC	2400
	ACGGGATGGT	GGGTCTGGGA	CCCCAATTCA	TTCTTATCTG	CCAAAGAATT	ATCTAGAAGC	2340

#### 15 (2) INFORMATION FOR SEQ ID NO: 12:

20

#### (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1131 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: double
  - (D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

25	CACTGCACCA GCTTTGTTAT CTGTAAAATG ATGATAATAC CAACACCTTC TTCTTGGGGT	60
	ACTGAAGATG AGAGAACATG ATATGTGTAA AGTGCCTTCC ACAATACCCA GAACATAGCA	120
20	AACATGTAAT GAATGTAGTA ATAGTAATTA TTTTATTTTC TTTTGATTCA GTTGGGACTA	180
30	TGTTCAGCTG TAACAGAATA CCCAAAATAA CTGTTTTAAA CAAATTAAAG TTTWGTTGTG	240
•	AAGTTTTGTT ACGAATTCAG ACAATCCAGG GCTTTTATAG ATGCACCAGG ATCAGCAGGT	300
. 35	ACAAAGGCAT CTTTCCTGAT TTCTGCCAGT CTCAATGCAT GGGTTGCAAT CCAGARTCCA	360
	RGATGGCAGT TCCAGCCCTG GTTĄCGCCCA TATTAGCACA CAGAAAGAAA GAGAAAGGGA	420
	TGTGCCTCTT CACTTTAATC ATAGCTCCCA CTAGATGCAC CCACTACTTC TGCTGATACT	480
40	CCATTAGCTA ATGCTTGCTT ACATGGTCAC ACTTAGTTTC CAGAGAGACA TGTCTGGACA	540
	GTCATGTGCT CAATTAATAT CCAAGTGTCC AATTACTGAG AAAAAAAGAA ACTAGCACCT	600
45	TTGCTTGGTT GCATTCCTCT TAGCATAAGC CACATTCTTT TTATGAAGTT GTCCTCAGTT	660
	ACTTGGATGC CTCAGTTGTC CTTTCAWITA GAAAWGCYCC TKGGACAYCC TGAAWCTGAC	720
	TTCTTTTGTC ATCAGCACCA TCACTACCAC TGCCYTCTTC AAAGCCACCA CGTTCTGTCC	780
50	CCAGGATGGT TGCAACAACC ACCATAGGGA CTTTTTGCCT TCTACTTCCA CACAATAGNC	840
	CAGAGTAAGC TTTTGAAAAT GTAGGTCAGA TCATGTCTCT CTCTTCCTCT TCAAAACCCT	900
55	CCCGATGGCT TTTCATATTA CTCAAAAGAA AACCTAAAAC TTTGCTGTGA GATCTATGTG	960
	ACCCGGCTTA TICTICCTCT TACTITATCT CTGTATTGCT CTTCCTCACT CTACTCCAGC	, 1020
60	CATCCCACCT CCTTGCTGCT TGTCCTATAC TCCTAAAAGA AGTTCAGTCT TCCCTTATGA	1080

(2) INFORMATION FOR SEQ ID NO: 13:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 941 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	60
(2) INFORMATION FOR SEQ ID NO: 13:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 941 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	. 60
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 941 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	. 60
(A) LENGTH: 941 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	. 60
10 (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	. 60
(D) TOPOLOGY: linear	. 60
	. 60
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:	, 60
15 GGCACGAGTA GCATTTCATT TAATCTGCAG GTATATTCTC CCAACAGTTT ATTGTCATGT	
GATGTCCTCA GCCAAGATTG TRAGGCAGAG AGGAGCTGTC CCAACCTACT ATACCACCGA	120
and the second s	180
TGTCCTCTGT AGCAAACCGG AAAAGTCAGT GACAGAAGAT GCCGCTAGCG GTTTGAGCCA	240
GAGAATGACA GCTCTGGTTT GGAGAAAAGG GCCGGATGGT GGCTCTAGAA AGCCCATCCT	300
TCTGCTCTTC TTTTTTCTCC CCCTTATATT GTGCTTTCAT TCATTCATTC ATTCATCAAA	360
CATTTGTTGA GCACCTATTA TGTGTCAAGC TCTGTGCTAG CCTCTGGAAA ACCTGCCCTC	420
30 ATGTAGCTCA CTGTGGAGTA GGAGAAACAA TGACTACACT ATGATAAGCA CGGGTTGTCA	480
GGGTCTCACA GAGCAGTGGC CCCTCATCCA GACCGATGAG GTCAAAGAAG GCATCCAGGC	540
GAGGATGGTG TCAGAGCTAA CTGAAGAATG AGAGGGAGCT GCACCASCAG GGGTTGGAAC	600
35 TGAAGGTGGC AGTGCCTGGA GTCTTGATTC CAGCAGAGGG AGAGCAGTCT GTGAAAAAGGC	660
ACCAAGGGTG GGAGAGGGCA GAGCACATGG AGGAACTTCA GGTAGTTCTG GATGGCSCTG	720
40 GGGCAAAGCT AGAGAGGTAA GAAGAATCTA CAAATGTTCC TCGAGTTACA TGAACTTCCA	780
TCCCAATAAA CCCATTGGAA ACGAAAAATT TAAGTCAGAA GTGCATTTAA GGCTGGTCCG	840
AGTAGAATGA TTTTTACAAC GAATTGATCA CAACCAGTTA CAGATGTCTT TGTTCCTTCT	•
45	941
CCACTCCCAC TGCTTCACCT GACTAGCCTT TAAAAAAAAAA	
,	
(2) INFORMATION FOR SEQ ID NO: 14:	
(i) SEQUENCE CHARACTERISTICS:	
(A) LENGTH: 843 base pairs (B) TYPE: nucleic acid	
(C) STRANDEDNESS: double (D) TOPOLOGY: linear	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

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	CNAGGGATAA CCCCAAAGNT GGGAAATAAA CCCTCAATTA AAGGGGGAAC CAAAAAGCTG	6 <u>0</u>
	GGAAGTTCCC CCCCGCGGTG GCGGCCNGNT CTAGGAACTA GTGGAATCCC CCGGGGCTGC	120
5	AGGGAATTCG GCACGGAGTG GGAATGTTGT TTGTATGATA CTATTTCCAC AAWATGCATT	180
	GAGACTTGGT KTGTGGCCTA GGACATGGTC AATTCTTTYT AAATATTCCG TGAATTTCTT	240
• •	TAGTGCATAT TCTCCGATGG GGGCTGTGGG GACAGAGTTC TAAATATGCC CATTAGATTA	300
10	AATCTCTTCA TTCTGTTGCT CACATCTTCT ATATCCTTAT TAATCTGTCA ATCTCTTCAA	360
	GAGAGGTGTT ATTAAAATCT CTCACTGTAT GTGTCACTTT GCCCTTAAAA TTCTGATGAT	420
15	TTGCTTTATA AATGGTTATA ACCATTTTCC AGGAAGAACA TTAAAGAACT TTCCATTGGC	480
	ATTATCCAGT TTCCCTCAAA ATACTGGTTT TTTTTATTTT GGCTNCTAAG CAGCTATGAA	540
20	TCCAGTTTCT CAGAAGCCCT TGTCTCAAGG CATTTGTTTC CAGATTACCT TGTTAGCATC	600
20	CACACTATGG GCTATTTTAG AAAAACAAAA AAAGTATCAA AATCATATAG CTATGATTTT	560
	CCTGTGCTTG AAGGAGCCTT AAAGCTCATC TAGTCCAGCC AGTATTTGTT CATCCAAATT	720
25	CTGCCAAGAA ATCTCTATTG TCAAGATATT CTTTACCATC TTTGGGACAT TCTCATTATT	780
	AGAAACAAAT CCTAAGAAGA AATTCTGCCA TAKACAACCC ATCCGTTCTT TAAAAAAAAA	340
30	AAA	343
50		
	(2) INFORMATION FOR SEQ ID NO: 15:	
35	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1018 base pairs (B) TYPE: nucleic acid	
40	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:	
	CTGTAATTIT TAATTITCAT ATACCGTGCT TTGATTCTAA TTTTATTTTT TGAGTTCTCT	60
45	GAAGGITACA TATACAGAGI GCTICAGGAA IGATCATITI GITATTATIC AIGCITCITA	120
	ACAATGTTGT TTTAGTCCAA GAAGATAATT GCCAGAGAAA GAATACAGTG CAGGAAAGAA	180
50	GARGCTGGAG CCAGTGGTGA AGARGGATTG AGARGACAGA CATTGTGGGA ATGAAATCAT	240
30	GANTANTOGT GITTITGANT TOTOCANANA CITCIACANA CONTGANATO TIGGAOTITA	300
	WHILE SALES OF THE	

AATCTAATTG TTGAAAAATT CCCCACATTC CTTGTATCCC TTAGGTTGAG CATAATTCCA

CATCCGTGGA CTGATGCACT TCCCAAGAGG GGGCCTCATT AACTCTTCCG AGGCAGCAGC

AGCAAGGGCA CCCCCTCCTT TCCCCCCACA CCCCAYTTCT CATGGCTCTT CTTTCTCTCA

TCTCATGCTT AGGTTAGAAA AGGGCACAAG GTAAGGAAGC CCTTGGGAAT AGGCTGAATC

	TGGCTATCTA ATTTGGTGCC AAATACTTAA TGTGCTTGAA TTTAAAAACA GCAAACATGT	600
5	AGAAAGGTAA TTATAATTAT GAGGCCAGTT CTTTAAGCTA GCTTTTTTTC CCCTCTCAAA	660
	CAGCATATTG GCTTGGATGT CAGCAGGAGA AAGTGTTTTT TGCAATACAC ATAATGCATA	720
	TATGGTCCTG TTAGCAATCT ATAGAAAATA GATATTGCTC ATTAAGGTAA ATATTTTTGT	780
10	TGATGAATGA TCTGGAATGG TCTGGACTTG TTGTGTGAAC AGGAAATTGC TCTGTAGGCT	840
	TTGACTTGTG AGGTAAAGAG TGAGGCTGGT AAGATTAATT AAAGTAAATA CTGTGACAAT	900
15	AGGATGTCAA AACCAAAAAC GTGTTTCTGA AACTCAAGGA ATTAATGACA CATAGGGAAG	960
13	TTTTTGCCAT ATTAAGCATA GAGTAGGAGA GGCAAGTCAA GAATAAAAAA AAAAAAAA	1018
20	(2) INFORMATION FOR SEQ ID NO: 16:	
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 661 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:	
30	TTTAAGAAAT TAGTGAATCC CCGGNTGCAG GGAATTCGGC ACGAGGAGGA GGCCGTCAGC	60
	TEGERAGAGE GEAGGATGGE AGETGYTECE CEGGGTTGEA CECECECAGY TETGETGGAE	120
35	ATAAGYTGGT TAACAGAGAG CCTGGGAGCT GGGCAGCCTG TACCTGTGGA GTGCCGGCAC	180
	CGCCTGGAGG TGGCTGGGCC AAGGAAGGGG CCTCTGAGCC CAGCATGGAT GCCTGCCTAT	240
40	GCCTGCCAGC GCCCTACGCC CCTCACACAC CACAACACTG GCCTMTCCGA GCTGCTGGAG	300
40	CATGGAGTGT GTGAGGAGGT GGAGAGAGTT CGGCGCTCAG AGAGGTACCA GACCATGAAG	360
	GTGCGCAGGG CAGGGCTCGG ACCTACCCCA GGAATGTCCT GCCCTGGGAA TGACAACACA	420
45	GTCCACACCA TGCACGGGA GGCAAACAGG GGCAGCTGAC CCAGCCCAGG GGTCAGANGA	480
	GGTCTTGCCG AGGAAGTGGC AGCTAAGCTG ATACCTGATA TGCACWAGKC AGCCARGYGG	540
50	AGACAGGCAA GGAAGAAGCT TGTTTTGAGG ACAGAATTTT CTAGATCACT CAGCACCATC	600
50	TGGCTTTTGG GGCTTTTGT TTTATTTTGT TTTTGAGACG GGGTCTCGCT CTGTCGCCCA	660
	N	661

(2) INFORMATION FOR SEQ ID NO: 17:

N

55

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(i) SEQUENCE CHARACTERISTICS:

	(A) LENGTH: 553 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:	
	GGCACAGGGC TATTTGCCCC TCTCTCCACA TGACAGAACT GCTCTAAGTT TCTTTGCTGC	• 60
10	TCTTCTCAGC TGTCAGACGG CTTGCTGCTT GTTTTCCACA CCACCATGTC TATTCTTTGC	120
	TGTCCTTWAC TCTGCCTGTT TTTTTCCTTT TGTATTTCTT CTGGCTCTTG TCCCTTTTCC	180
1.5	CACGTGTCWC AGCTTTCCTT TATTGCCACT TTCAGTCAGA GCAGTCCTGT GCTTCTGGTG	240
15	CCGGCATACA ATACTTACTT GAGTTTCTTG GCTTTTCTTG ACTGTGCATC TCTTACTTCA	300
	ACATAGGAAT AGCCTGTCAT AGAATTTCTC CAGTTCCAGG GCTCAAGAGG GAGAGTGCCA	360
20	GAAAATTGAG ACTGTTTTCC CTGTCTTGGA TTGAATTCAT AAAGCAAAAC CAGTGTTTGT	420
*	GTGAGGGTTT GCTGTGTCAT GCCTATAGGT TGTTTGGGTG CAAACCTATA GAATCCAGCC	480
05	TGCGAAAAGA AAGRAACCAG AGAATANCAG CATCAGAACA ATGCTTGACA TCATTTCTCA	540
25	ATCAAGCAGT CCA	553
30	(2) INFORMATION FOR SEQ ID NO: 18:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 869 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:	
	GGCACGAGCT GCCAACACTG AGGTCTTCGT GGCTTCTCAC ATCTAGATGT ATCCCTCTCA	6
•	AATCTATCCT CTATCCAGGC ACCAGATTGA GGTATCTAAA ATGTCAACTT TCCAGTTACT	12
45	CCTTCTTATA CTAGCCCAAT CAACTTACAA GATAAAGTCC AAGCCCCTTC ATATGACAAA	18
	CCACACCCTG CTTAACTCTC CAGGTTTGAA TCCTTCATCT CCTACTTTAA ACTTTAAAAC	24
50	CCAGCAGCAC GAAAGTGTCT CCTATGCATG TTGCCATATG CGTTCTCTCC ATCATGCATT	30
	TGCCTGAGCA AGATGTCTTG AGTTAACATC TTATTCTTTA AGACTCATTG TGGTGGTAGA	36
	CAGCCTTTAA TAACGGATCC TTGGCCAGGC ACAGTGACTC ACACCTGTAA TCCCAGAACT	42
55		48
	AGAGAGATAT CCCATCTGTA CCAAAAATTT AAAAAAATAT TAGCAGGGAG TAGTGGCATG	54

CACAAGTGGT CCCAGCTCCA TGGGAGASTG AGGTAGGAAC ATCACTTGAG CCCAGGAAGT

60

	CAAGGCTGCA GTGAACCATG ATCAGAACAT TGCANTCCAG CTTGGGTAAC AGAGTGAGAC	660
	CTTAGGTCAG AAAAATGAAT AAATAAGCAT AAAATTTTAA AAACTTAGCC AGGCATGGTG	720
5	GCACACATCT GTGGTCCCTG CTACTTAGGA GGCTGAGGTG AGAGGATCCT TGAGCCCAGG	780
	AGGTCAACAC TACAGTGAGC TATGATTGTG CCACTAAACT CCAACCTGGG TGAAAAAGCA	840
	AAACCCTGCC AAAAAAAAA AAAAAAACT	· 869
10		
	(2) INFORMATION FOR SEQ ID NO: 19:	
15	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 959 base pairs (B) TYPE: nucleic acid	
20	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:	
	GGCGAGCCGA GATCGTGCCA TTGCACTCCA GCCTGGGCAA CAAGAGTGAA ACTCTGTCTC	60
25	AAAAAAAAA AATTATAATA CTATATGCCA TAAAATGACA TTTCATATTT AAAGAGTTTT	120
ě	TTAAAACTCT TGTATTCACA TGCCATAATT TGAAACCCTA TTTCACTGAA TGAGAATGGT	180
30	ATCIGITGIC CICATITITI CATITITATC CITAACAATT TCCACCACAG CCAGIGCATA	240
	TAATGGCAAT GACACCCAGG GATGGAATGA TAAGTTCCAT CRCMGCTCAG TCAAGACGCA	300
25	GACTTGATGT GGCCCCAACA ACAGTCAATA ATGGAGTCTC CAAAATAAAG CTCTATAGGA	360
35	AAGGTAAATA CCCGCTGCAC AAGAAACCAC AGCATCTAGG TTCTAACCCC ATCTCTATGA	420
	AGAGCTTGCT GGGAGAGTTT TGACATTWAA CAATCTGTCT GATKGCCAAT TTTYTTCTTC	480
40	TATAAAATGA TAATGTTKGA YTCAAAGATC CAAAGTCAAT TCATGGTCTA AAACTTAATG	540
	ATTTTTTTAG GTTTTGKGAC ATTTCACTGT ACACTGTAGT AATTTATATC TTATTTTCCC	600
15	ACTAATTTAG AAAAATATYT AAATGATCCT TAATTGGCAA TGGGTCCTAA GAATTTTGTT	660
45	TTAAATCCCT GTTACCCAAA AGAGCCCTTT TTTGTATCTC GCAGTAGTTA CAAGGATCTT	720
	TCTAAATCTT AAAAAAAAA AAAAAAGAAA GAAAGAAAAG AAAAGAAAAA AAGTCAGCCG	780
50	GGCGTGGTGG CTCATGCCTG TAATCCCAGC ACTTTGGGAC CAAGGTGGAC AGATCACGAG	840
	GTCAGGAGAT GGAGACCATC CCGGCCAACA TGGAGAAACC CTGTCTCTAC TAAAAAAAAAA	900
55	AAAAACTCGA GGGGGGCCCG GTACCCAATN CGCCGGCTAG TGGTCGTAAA ACAATCAAA	959

<sup>(2)</sup> INFORMATION FOR SEQ ID NO: 20:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1446 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

10	CGGGGCAGGG CTGTGTGGCA CCGCCAGGGA GCGGGCCCAC CTGAGTCACT TTATTGGTT	- 60
10	CAGTCAACAC TTTCTTGCTC CCTGTTTTCT CTTCTGTGGG ATGATCTCAG ATGCAGGGGC	120
	TGGTTTTGGG GTTTTCCTGC TTGTGCCAAG GGCTGGACAC TGCTGGGGGG CTGGAAAGCC	180
15	CCTCCCTTCC TGTCCTTCTG TGGCCTCCAT CCCCTCATGG GTGCTGCCAT CCTTCCTGGA	240
	GAGAGGGAGG TGAAAGCTGG TGTGAGCCCA GTGGGTTCCC GCCCACTCAC CCAGGAGCTG	300
20	GCTGGGCCAG GACCGGGAGA GGGAGCACTG CTGCCCTCCT GGCCCTGCTC CTTCCGCAGT	360
20	TAGGGGTGGA CCGAGCCTCG CTTTCCCCAC TGTTCTGGAG GGAAGGGGAA GGAGGGGGTC	420
	TTCAGGCTGG AGCCAGGCTG GGGGTGCTGG GTGGAGAGAT GAGATTTAGG GGGTGCCTCA	480
25	TGGGGTGGGC AGGCCTGGGG TGAAATRAGA AAGGCCCAGA ACGTGCAGGT CTGCGGAGGG	540
	GAAGTGTCCT GAGTGAAGGA GGGGACCCCC ATCCTGGGGG ATGCTGGGAG TGAGTGAGTG	600
30	AGATGCCTGA GTGAGGGTTA TGGGGAGCCT GAGGTTTTAT GGGCCTGTGT ATCCCCTTCT	660
30	CCCGGCCCCA GCCTGCCTCC CTCCTGCCCG CCTGGCCCAC AGGTCTCCCT CTGGTCCCTG	720
	TCCCTCTGGT GGTTGGGGAT GGAGCGGCAG CAAGGGGTGT AATGGGGCTG GGTTCTGTCT	780
35	TCTACAGGCC ACCCCGAGGT CCTCAGTGGT TGCCTGGGGA GCCGGACGGG GCTCCTGAGG	840
	GGTACAGGTT GGGTGGGCCC TCCCTGAGGG TCTGGGGTCA GGCTTTGGCT CTGCTGCCTC	900
40	TCAGTCACCA AGTCACCTCC CTCTGAAAAT CCAGTCCCTT CTTTGGATGT CCTTGTGAGT	960
40	CACTCTGGGC CTGGCTGTCG TCCCTCCTCA GCTTCTTGTT CCTGGGACAA GGGTCAAGCC	1020
	AGGATGGGCC CAGGCCTGGG ATCCCCCACC CCAGGACCCC CAGGCCCCCT CCCCTGCTGC	1080
45	TTTGCGGGGG GCAGGGCAGA AATGGACTCC TTTTGGGTCC CCGAGGTGGG GTCCCCTCCC	1140
	AGCCCTGCAT CCTCCGTGCC STAGACCTGC TCCCCAGAGG AGGGGCCTTG ACCCACAGGA	1200
50	CGTGTGGTGG CGCCTGGCAC TCAGGGACCC CCAGCTGCCC CAGCCCTGGT CTCTGGCGCA	1260
50	TCTCTTCCCT CTTGTCCCGA AGATCTGCGC CTCTAGTGCC TTTTGAGGGG TTCCCATCAT	1320
	CCCTCCCTGA TATTGTATTG AAAATATTAT GCACACTGTT CATGCTTCTA CTAATCAATA	1380
55	AACGCTTTAT TTAAAGCCAA AAAAAAAAA AAAAAACTCG AGGGGGGCC CGTACCCAAT	1440
	TCGCCA	1446

(2)	INFORMATION	FOR	SEQ	ID	NO:	21:
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1471 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

	(D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:	•
	CAAAAAATAA TAATGATAAT TTAAAATAAA TAAGTAACTA ATAAAAAGAT TTTATATCCC	60
15	AGTCTTATGA TGTTGGTTGG CAAGGCTAGA TAAAAAGATG TTAGAATGAA AGAACATATT	120
13	TTTAGTGATA TGTAAATGAA GGATTCTACA ATAGTCATAT ATTTTTATAT GAATGAATGT	180
	TGGGTTGGGC TGGAGAGGTA TGTGTGTGTA AATATAAAGG TCTCACATTC AGAGTATAGC	240
20	TCTGAAATAA TGGAACTCAT GTCTACAATT CAACATGCAT CTGTATAGTT ACATCTCATG	300
	TAAATATACA CAGACATATT TTGCAGCCAG TAATTGACAG TTAATGTCCA AAACAGGTGA	360
25	TTGATAGGTA ACAGAAATTA GATAACCACC AATTTTGCCC AAGAGAAAGA CTAGAAGGAC	420
23	TAAAAGCAGT TGAATGTATG GTACTGACAT TGTCATAAGC AGTCTGATAA CCAGTTTATT	480
	GAAACGTGTG CATTAACAGA GAATTTAATT TTAAACCCAT AATTTCTCCT ATCCATTAAA	540
30	ATATTATAAT TGTTAGTAGT ATGAAACCAA CAGGAAATGT TTTTTAATCA TTTAGTGAGG	600
	TGATTCATTT GTTTCATGGG CAAACACTAT CCAGGAAAAG CCTTGCTTGC CTGTTTCCCA	660
35	AAGAGCTCTA AGAAATAGAA TCAAGTGTAA AATGGTTCAG ACCATTCAGG ATTTCTTGTC	720
75	ACTOTTOTCA ACCCCGATCT TCCTGTTATT ACTGATGTTT GAAACCCTGT CATTAGCCCC	780
	GGCCTGGTTA AAGCCCCTCA GAGTCACCTC TCATTCATAG CAATAGAATT CAACCCCAAG	840
40	TOGTTGATGG TGTCCCCAGC ACAGCCGAGA GACCTGATCT CTGGATTCAG TGCTTTTAGC	900
	TCTTCGAGTT TACCCTAAGA TACCTTCGGG CAATATTTTT AACCAACCCA AAAGCTCTTC	960
45	AGGTCATTTC TGAAGAGGAC AAGGTGAATC TTGGCTTGGA ACACCATTTT TGGGCTCTTG	1020
73	CTACTGAATG AATCAGAAAG GAATTTTTTC TGAAGAGCAT TAGAAAGTAA AGGAGATGTT	1080
	AAAATAAGTT CTTGAAGTAT GTTTTATATT TATCTAAAAC ACTGATTTTA AAAGTTTACA	1140
50	TTCAAATGTG TATTCAAAAG AAGTACTGAT TTGTAATTAT TATAGTTTGT GTGTATCATC	1200
	CCCTTTTAAC CGTGCCTAAC AACTGTACTT AAATTTTGTT TTCCTAGTGT AACAAATGTT	126
55	TCCCATAAGA TTTTCTAGAG CCAAATAATG GGAGTGAAAA ATTCCTTAAG TGTTATATAA	132
<i>JJ</i>	GAAAATATAT TAGAAAATCA GCTTTGGATT ATACGATTTC TAAAATATAC TAATACAGAA	138
	TCCTCAGTAA TATGTTTTGA ATTGGATTTT TTCTCAGAAC TGTTACATAA TAAATAATAC	144
60	ATCAACCAGA AAAAAAAAA AAAAAAATTN C	147

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5	(2)	INFORMATION	FOR	SEQ	ID	NO:	22:
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1402 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:	
15	AGGGACGTCT TGCCTGAGGA GATGCCCATT TCTGTCCTGG RTTACCCTCA CTGCGTGGTG	60
	CATGAGCTGC CAGAGCTGAC GGCGGAGAGT TTGGAAGCAG GTGACAGTAA CCAATTTTGC	120
20	TGGAGGAACC TCTTTTCTTG TATCAATCTG CTTCGGATCT TGAACAAGCT GACAAAGTGG	180
20	AAGCATTCAA GGACAATGAT GCTGGTGGTG TTCAAGTCAG CCCCCATCTT GAAGCGGGCC	240
	CTAAAGGTGA AACAAGCCAT GATGCAGCTC TATGTGCTGA AGCTGCTCAA GGTACAGACC	300
25	AAATACTTGG GGCGGCAGTG GCGAAAGAGC AACATGAAGA CCATGTCTGC CATCTACCAG	. 360
	AAGGTGCGGC ATCGGCTGAA CGACGACTGG GCATACGGCA ATGATCTTGA TGCCCGGCCT	420
30	TGGGACTTCC AGGCAGAGGA GTGTGCCCTT CGTGCCAACA TTGAACGCTT CAACGCCCGG	480
30	CGCTATGACC GGGCCCACAG CAACCCTGAC TTCCTGCCAG TGGACAACTG CCTGCAGAGT	540
	GTCCTGGGCC AACGGGTGGA CCTCCCTGAG GACTTTCAGA TGAACTATGA CCTCTGGTTA	600
35	GAAAGGGAGG TCTTCTCCAA GCCCATTTCC TGGGAAGAGC TGCTGCAGTG AGGCTGTTGG	660
	TTAGGGGACT GAAATGGAGA GAAAAGATGA TCTGAAGGTA CCTGTGGGAC TGTCCTAGTT	720
40	CATTGCTGCA GTGCTCCCAT CCCCCACCAG GTGGCAGCAC AGCCCCACTG TGTCTTCCGC	780
40	AGTCTGTCCT GGGCTTGGGT GAGCCCAGCT TGACCTCCCC TTGGTTCCCA GGGTCCTGCT	840
	CCGAAGCAGT CATCTCTGCC TGAGATCCAT TCTTCCTTTA MTTCCCCCAM CCTCCTCTCT	900
45	TGGATATGGT TGGTTTTGGC TCATTTCACA ATCAGCCCAA GGYTGGGAAA GCTGGAATGG	960
	GATGGGAACC CCTCCGCCGT GCATCTRAAT TTCAGGGGTC ATGCTGATGC CTCTCGAGAC	1020
50	ATACAAATCC TTGCCTTTGT CAGCTTGCAA AGGAGGAGAG TTTAGGATTA GGGCCAGGGC	1080
50	CAGAAAGTCG GTATCTTGGT TGTGCTCTGG GGTGGGGGTG GGGTGTTTCT GATGTTATTC	1140
	CAGCCTCCTG CTACATTATA TCCAGAAGTA ATTGCGGAGG CTCCTTCAGC TGCCTCAGCA	1200
55	CTTTGATTTT GGACAGGGAC AAGGTAGGAA GAGAAGCTTC CCTTAACCAG AGGGGCCATT	1260
	TTTCCTTTTG GCTTTCGAGG GCCTGTAAAT ATCTATATAT AATTCTGTGT GTATTCTGTG	1320
	TCATGTTGGG GTTTTTAATG TGATTGTGTA TTCTGTTTAC ATTAAAAAGA AGCAAAAATA	1386

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(2) INFORMATION FOR SEQ ID NO: 23:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1047 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:	٥
13	GGCACAGGGG ACTACAGGCA CCCACGACCA TACCCAGCTA ATTTTTGTAT TTTTTTTTAG	60
	AGATGGGGTT TCACGATGTC GCCCAGGCTG GTCTTGAACT CCTGGGCTTG AGCGATCTTC	120
20	CCATCTTTCC ATCTTGGCCT CCTAAAGTGC TGGGACTGCA GGCATGAGCC ACCATGCCCA	180
	GCCAAGATTC TTATTGATTA CCATGTTGCT TCAAGAAGCC AAGCCAGTTT CCAATATTCC	240
25	CCATTTGCTG GAGTCTTGGT ACTTTGGGTA GAAGCAACTG GTAAATTGTT AATTGGAACA	300
23	NTTGGTGGTG TAGATAACCA CGTATGGCCA AACCTAGAGC ATCTAGGCTC ACAATTACTA	360
	TCCTGACTTG ATAACAAGTG TTCTGATATT AACCTGAAAA TGGGAATAAT GCCAAATCTG	420
30	TGTAACTTAA CATCTATATA CACAGTGGGG AGAACTGAAG TTATTAAACC TGGAATCTCT	480
	GIGATCAAGG CTAACAGTAG TTATCTAAGA AGCAAAGGAC CTACAATTCT TAGACTTGGA	540
35	GTCATATTCT TTAAGGACGT GTTCTGAAAC TATATCAAGC ATCTGGTTTC CACGTATTTC	600
33	TCCCTCAGAA ATTATGAAGT ACAAGTAAAA ATGAAGGTAC AGGGTAAGAC ACATGCTGCT	660
	TTCTTGCTCT TGAGTGGAGA CAGTTTTCCA GCCATCTTAA CCCCTTWACA CAAAACAATT	720
40	TGTGTTTTAT AGCAAATAAG TGACTCAACA TAATTTCAAT ATGATGTTTA TCCACCAGTA	780
	CTTTCCTTTC AGCTTCTAGT CCCATAARTG GTTTGTGAAG TCATCGGTTA CATTAGCCAA	840
45	GATAGGCCTA GACTTGAAGT CTAGAATGTT TTTCCCACTA TATGCCAAAG TAGAATGTGG	900
43	GTATCTCAGG GTCATTTTTG TTGTTCAATT TCCCACCTGT ACAGTTGTTA TGATTCACTT	960
	TCCTTATGTG TCTAATAAAT CTTGTTCCAT GAAATGATCA AAAAAAAAA AAAAAAAACT	1020
50	CGAGGGGGG CCCGGTACCC AAATCGC	1047

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(2) INFORMATION FOR SEQ ID NO: 24:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 990 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

#### (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2	(xi)	SECUENCE	DESCRIPTION:	SEO	ID NO:	24:
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5	TTGGAAAGGG	TCTAGCTCTT	TCTCATTCAC	CAACTATATT	AGAAGCACTT	GAGGGAAATT	60
	TACCACTCCA	AATCCAAAGC	AATGAACAGT	CTTTTCTGGA	TGATTTTATT	GCCTGTGTCC	120
10	CAGGATCAAG	TGGTGGAAGG	CTTGCAAGGT	GGCTTCAGCC	AGATTCATAT	GCGGATCCTC	. 180
10	AGAAAACATC	TTTGATCCTG	GAATAAGGAT	GATATTCGTT	GTGGTTGGCC	TACCACCATA	240
	ACTGTTCAAA	CAAAAGACCA	GTATGGGGAT	GTGGTACATG	TTCCCAATAT	GAAGGTAATT	300
15	ATAACTGGAT	TAAATTAGCA	GACATCTATA	TACTGGCTGC	AATGACTGAT	AAAATTTTAG	360
	AAATGCCAAG	TGCTGAGRGT	CCATTTGTTC	TACCCTCTTT	ATATAAAGGG	TGATGCTGAA	420
20	AGTTTGTTTA	AATGACTTGT	TTATATTAAT	TAGTCCCCAA	GTGTCCAAGT	TACACCTGTT	480
	TTTTTTTTGTGA	GTTTGTTCTT	TACATTTTGC	TACCTGTTAC	GGGGACTCAA	AGGAGGGATA	540
	AGAAAGTATC	CATCTAAAGA	GTGCTAGACA	CATACAGTGA	AGCCCCTCAA	TATGTATTGA	. 600
25	TTGAATAAAT	GCATGAAAGA	ATACATTTTT	AAATTTTGTG	TATAGTTTTG	AAAGACTCAA	660
	GTACGTTCTG	TGTTTGGTAT	TACTGAAACC	ACATTTTAAA	AATAACACTC	ATTAAGTTAG	720
30	AAATATATGA	GTTTAGATTG	TAAAAGAATG	AGGAATTGAA	ATAGTTGTAT	ACCATATTGA	780
50	TGAATATAGA	GTTTTTAGGA	TACCTCTTAC	CTGAAATATT	AATAATAATG	TTTNCAGAGC	840
	ATATTATACA	TAATTATTTG	TGATTTAATC	TGTTAATATG	AATATCTCAT	TTAAAACTTT	900
35	TATTTCTGAA	AAAATTATAT	TGAATAAAAT	TTTATATAGG	CAGTCCCCAG	CCCTTTCCTC	960
	CTTCAAAGTT	GTCTTATAGA	GTGATTGGTT		•		990
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#### (2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1208 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:

			•				
	TAATCGCTAC	TATAGGGAAA	GCTGGTCGCT	GCAGGTACCG	GTCCGGAATT	CCGGGTCGAC	60
	CCACGCGTCC	GAGCGAAATG	GCGCCTCCGG	ccccccccc	GCCTCCGC	GCTCCGGGG	120
55	AGGTAGACGA	GCTGTTCGAC	GTAAAGAACG	CCTTCTACAT	CGGCAGCTAC	CAGCAGTGCA	180
	TAAACGAGGC	GCASGGGTGA	AGCTRTCAAG	CCCAGAGAGA	GACGTGGAGA	GGGACGTCTT	240
60	CCTGTATAGA	GCGTACCTGG	CGCAGAGGAA	GTTCGGTGTG	GTCCTGGATG	AGATCAAGCC	300

•	CTCCTCGGCC	CCTGAGCTCC	AGGCCGTGCG	CATGTTTGCT	GACTACCTCG	CCCACGAGAG	360
5	TCGGAGGGAC	AGCATCGTGG	CCGAGCTGGA	CCGAGAGATG	AGCAGGAGCK	TGGACGTGAC	420
3	CAACACCACC	TTCCTGCTCA	TGGCCGCCTC	CATCTATCTC	CACGACCAGA	ACCCGGATGC	480
	CGCCCTGCGT	GCGCTGCACC	AGGGGGACAG	CCTGGAGTGC	ACAGCCATGA	CAGTGCAGAT	540
10	CCTGCTGAAG	CTGGACCGCC	TGGACCTCGC	CCGGAAGGAG	CTGAAGAGAA	TGCAGGACCT	600
	GGACGAGGAT	GCCACCCTCA	CCCAGCTCGC	CACTGCCTGG	GTCAGCCTGG	CCACGGGTGG	660
15	TGAGAAGCTG	CAGGATGCCT	ACTACATCTT	CCAGGAGATG	GCTGACAAGT	GCTCGCCCAC	720
13	CCTGCTGCTG	CTCAATGGGC	AGGCGGCCTG	CCACATGGCC	CAGGGCCGCT	GGGAGGCCGC	780
	TGAGGGCCTG	CTGCAGGAGG	CGCTAGACAA	GGATAGTGGC	TACCCRGAGA	CGCTGGTCAA	840
20	CCTCATCGTC	CTGTCCCAGC	ACCTKGGCAA	GCCCCTGAG	GTGACAAACC	GATACCTGTC	900
	CCAGCTGAAG	GATGCCCACA	GGTCCCATCC	CTTCATCAAG	GAGTACCAGG	CCAAGGAGAA	960
25	CGACTTTGAC	AGGCTGGTGC	TACAGTACGC	TCCCAGCGCT	GAGGCTGGCC	CAGAGCTGTC	1020
23	AGGACCATGA	AGCCAGGACA	GAGGCCAGGA	GCCAGCCCTG	CAGCCCTCCC	CACCCGGCAT	1080
	CCACCTGCAT	CCCTCTGGGG	CAGGAGCCCA	CCCCCAGCAC	CCCCATCTGT	TAATAATAT	1140
30	CTCAACTCCA	RGGTGTTCCA	CCTGAAAAAA	AAAAAAAA	AAAAAAAAA	AAAAAAAA	1200
	ААААААА						1208

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#### (2) INFORMATION FOR SEQ ID NO: 26:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1922 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:

GTGCTGCGCT	ACTGAGCAGC	GCCATGGAGG	ACTCTGAAGC	ACTGGGCTTC	GAACACATGG	60
GCCTCGATCC	CCGGCTCCTT	CAGGCTGTCA	CCGATCTGGG	CTGGTCGCGA	CCTACGCTGA	120
TCCAGGAGAA	GGCCATCCCA	CTGGCCCTAG	AAGGGAAGGA	CCTCCTGGCT	CGGCCCCGCA	180
CGGGCTCCGG	GAAGACGGCC	GCTTATGCTA	TTCCGATGCT	GCAGCTGTTG	CTCCATAGGA	240
AGGCGACAGG	TCCGGTGGTA	GAACAGGCAG	TGAGAGGCCT	TGTTCTTGTT	CCTACCAAGG	300
AGCTGGCACG	GCAAGCACAG	TCCATGATTC	AGCAGCTGGC	TACCTACTGT	GCTCGGGATG	360
TCCGAGTGGC	CAATGTCTCA	GCTGCTGAAG	ACTCAGTCTC	TCAGAGAGCT	GTGCTGATGG	420

•	AGAAGCCAGA	TGTGGTAGTA	GGGACCCCAT	CTCGCATATT	AAGCCACTTG	CAGCAAGACA	480
	GCCTGAAACT	TCGTGACTCC	CTGGAGCTTT	TGGTGGTGGA	CGAAGCTGAC	CTTCTTTTTT	540
5	CCTTTGGCTT	TGAAGAAGAG	CTCAAGAGTC	TCCTCTGTCA	CTTGCCCCGG	ATTTACCAGG	6.00
	CTTTTCTCAT	GTCAGCTACT	TTTAACGAGG	ACGTACAAGC	ACTCAAGGAG	CTGATATTAC	660
10	ATAACCCGGT	TACCCTTAAG	TTACAGGAGT	CCCAGCTGCC	TGGGCCAGAC	CAGTTACAGC	. 720
10	AGTTTCAGGT	GGTCTGTGAG	ACTGAGGAAG	ACAAATTCCT	CCTGCTGTAT	GCCCTGCTCA	780
•	AGCTGTCATT	GATTCGGGGC	AAGTCTCTGC	TCTTTGTCAA	CACTCTAGAA	CGGAGTTACC	840
15	GGCTACGCCT	GTTCTTGGAA	CAGTTCAGCA	TCCCCACCTG	TGTGCTCAAT	GGAGAGCTTC	900
	CACTGCGCTC	CAGGTGCCAC	ATCATCTCAC	AGTTCAACCA	AGGCTTCTAC	GACTGTGTCA	960
20	TAGCAACTGA	TGCTGAAGTC	CTGGGGGCCC	CAGTCAAGGG	CAAGCGTCGG	GGCCGAGGGC	1020
20	CNAAAGGGGA	CAAGGCCTCT	GATCCGGAAG	CAGGTGTGGC	CCGGGGCATA	GACTTCCACC	1080
	ATGTGTCTGC	TGTGCTCAAC	TTTGATCTTC	CCCCAACCCC	TGAGGCCTAC	ATCCATCGAG	1140
25	CTGGCAGGAC	AGCACGCGCT	AACAACCCAG	GCATAGTCTT	AACCTTTGTG	CTTCCCACGG	1200
	AGCAGTTCCA	CTTAGGCAAG	ATTGAGGAGC	TTCTCAGTGG	AGAGAACAGG	GGCCCCATTC	1260
30	TGCTCCCCTA	CCAGTTCCGG	ATGGAGGAGA	TCGAGGGCTT	CCGCTATCGC	TGCAGGGATG	1320
50	CCATGCGCTC	AGTGACTAAG	CAGGCCATTC	GGGAGGCAAG	ATTGAAGGAG	ATCAAGGAAG	1380
	AGCTTCTGCA	TTCTGAGAAG	CTTAAGACAT	ACTITGAAGA	CAACCCTAGG	GACCTCCAGC	1440
35	TGCTGCGGCA	TGACCTACCT	TTGCACCCCC	CAGTGGTGAA	GCCCCACCTG	GGCCATGTTC	1500
	CTGACTACCT	GGTTCCTCCT	GCTCTCCGTC	GCCTGGTRCG	CCCTCACAAG	AAGCGGAAGA	1560
40	AGCTGTCTTC	CTCTTGTAGG	AAGGCCAAGA	GAGCAAAGTC	CCAGAACCCA	CTGCGCAGCT	1620
-10	TCAAGCACAA	AGGAAAGAAA	TTCAGACCCA	CAGCCAAGCC	CTCCTGAGGI	TGTTGGGCCT	1680
	CTCTGGAGCT	GAGCACATTG	TGGAGCACAC	GCTTACACCC	TTCGTGGACA	GGCGAGGCTC	1740
45	TGGTGCTTAC	TGCACAGCCT	GAACAGACAC	TTCTGGGGC	GGCAGTGCTC	GGCCCTTTAG	1800
	CTCCTTGGCA	CTTCCAAGCT	GCATCTTG	CCCTTGACA	A CAGAATAAAA	ATTTTAGCTG	1860
50	CCCCAAAAAA	AAAAAAAAA	AAAAAAACTY	CAGGGGGGG	CCCTACCCA	TTCGCCCTAT	1920
JU	AA						192

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(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1951 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

5		
3	TCGTCCCCAG AGCGGGCTGA GCCCCAGGCG SAGGGTGGCG GGGGAGCCTG GGGGAGCCGC	60
	CGCCACCTCC ACGGGCCTCT CTGAGCTCGG ACACCAGCGC CCTGTCCTAT GACTCTGTCA	120
10	AGTACACGCT GGTGGTAGAT GAGCATGCAC AGCTGGAGCT GGTGAGCCTG CGCCGTGCTT	180
	CGGAGACTAC AGTGACGAGA GTGACTCTGC CACCGTCTAT GACAACTGTG CCTCCGTCTC	240
1.5	CTCGCCCTAT GAGTCGGCCA TCGGAGAGGA ATATGAGGAG GCCCCGCGGC CCCAGCCCCC	300
15	TGCCTGCCTC TCCGAGGAAC TCCACGCCTG ATGAACCCGA CGTCCATTTC TCCAAGAAAT	360
	TCCTGAACGT YTTCATGAGT GGCCGCTCCC GCTCCTCCAG TGCTGAGTCC TTCGGGCTGT	420
20	TCTCCTGCAT CATCAACGGG GAGGAGCAGG AGCAGACCCA CCGGGCCATA TTCAGGTTTG	480
	TGCCTCGACA CGAAGACGAA CTTGAGCTGG AAGTGGATGA CCCTCTGCTA GTGGAGCTCC	540
25	AGGCTGAAGA CTACTGGTAC GAGGCCTACA ACATGCGCAC TGGTGCCCGG GGTGTCTTTC	600
23	CTGCCTATTA CGCCATCGAG GTCACCAAGG AGCCCGAGCA CATGGCAGCC CTGGCCAAAA	660
	ACAGTGACTG GGTGGACCAG TTCCGGGTGA AGTTCCTGGG CTCAGTCCAG GTTCCCTATC	720
30	ACAAGGCCAA TGACGTCCTC TGTGCTGCTA TGCAAAAGAT TGCCACCACC CGCCGGCTCA	780
	CCGTGCACTT TAACCCGCCC TCCAGCTGTG TCCTGGAGAT CAGCGTGCGG GGTGTGAAGA	840
35	TAGGCCTCAA GGCCGATGAC TCCCAGGAGG CCAAGGGGAA TAAATGTAGC CACTTTTTCC	900
JJ	AGTTAAAAAA CATCTCTTTC TGCGGATATC ATCCAAAGAA CAACAAGTAC TTTGGGTTCA	960
	TCACCAAGCA CCCCGCCGAC CACCGGTTTG CCTGCCACGT CTTTGTGTCT GAAGACTCCA	1020
40	CCAAAGCCCT GGCAGAGTCC GTGGGGAGAG CATTCCAGCA GTTCTACAAG CAGTTTGTGG	1080
	AGTACACCTG CCCCACAGAA GATATCTACC TGGAGTAGCT GTGCAGCCCC GCCCTCTGCG	1140
45	TCCCCCAGCC CTCAGGCCAG TGCCAGGACA GCTGGCTGCT GACAGGATGT GGCACTGCTT	1200
<del>-1</del> 2	GAGGAGGGC ACCTGCCACC GCCAGAGGAC AAGGAAGTGG GGCGCTGGCC CAGGGTAGGG	1260
	GACGGTGGGG CAATGGGGAG AGGCAAATGC AGTTTATTGT AATATATGGG ATTAGATTCA	1320
50	TCTATGGAGG GCAGAGTGGG CTGCCTGGGG ATTGGGAGGG ACAGGGCTTG GGGAGCAGGT	1380
	CTCTGGCAGA GAAGGATGTC CGTTCCAGGA GCACACGGCC CTGCCCCATC CTGGGCCTTA	. 1440
. 55	CCTCCCCTGC CAGGGCTCGG GCGCTGTGGC TCCTGCCTTG ATGAAGCCCG TGTCCTGCCT	1500
	TGATGAAGCC TGTGCCACCT GCAAGTGCCC GCCCTGCCCC TGCCCCAACC CCCACCGAAG	1560
	AGCCCTGAGC TCAGGCTGAG CCCAGCCACC TCCCAAGGAC TTTCCAGTGA GGAAATGGCA	1620
60	ACACGTGGAG GTGAAGTCCC TGTTCTCAGC TCCGTCATCT GCGGGGCTTC TGGGTGGCTC	1680

	CTGCCACTGA	CCTCACCGGC	ATGCTGGCCT	GTGGCAGGCC	TAGGACCTCA	GGCGGGGAGG	1740
5	AGGAGCTGCC	GCAAGGCCCT	GTCCCAGCAG	AAGAGGGAGG	CTTCCTGACT	GACACAGGCC	1800
3	AGCCCCATCT	TGGTCCTGTC	ACCCTGGCCC	CAACTATTAA	AGTGCCATTT	CCTGTCAAAA	1860
	ААААААААА	AAAATCGGGG	GGGCCCGGA	ANCCAATITC	CCCCAAAAAG	GGGGGTTATA	1920
10	AAAATTCCCN	GGCNGTGTTT	TTAAAAATTC	G			1951

#### 15 (2) INFORMATION FOR SEQ ID NO: 28:

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#### (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3989 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

#### (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

25	GGCACAGGCC GC	AGGGNACC	TATGGGCGCA	TATAGGTTGT	AATGAAACTG	TAGTCTCAGT	60
	TGGAAGCCTA GAG	CATGAAAT	GGGTCAGTGA	GCAAGGCTCT	ATTCCTAGTC	TCCAGCCATG	120
20	CCTGTGGAAC CT	GARCCCRC	TCTCAGCACA	TTGGACCCAG	GCAGATGYAA	AAAATTCACA	180
30	GAACTATGAT TT	GGACTCAA	GGGTTTGTAG	ATTTCCTCCT	TCATTCTAAT	TTCAGTGTCT	240
	AAAATTCTTG CA	TCCRTGAA	CGAGCTGGGC	ATTTGATGAG	ACAGGGCYGA	ATACTGCAGT	300
35	TTTCCTCCTA GA	AATCATCT	GGGGCATTTT	CTTTGAACTG	ATGGGAACAA	TAAGGCATAA	360
	CTGTTTGCAC AA	ACTTGGGA	TAARTGATTT	TGGGATAACG	ATCTACCAGA	ATGGGGATAT	420
40	TTCACCCTTG GT	TCTGAGAT	GCAAACCAAA	GAATATCATG	ACCAGCTTTC	AGGCCTCCTG	480
40	AAGTATATCT CT	CACATTGT	CCTGTTCTCA	TGCTGAGGAG	CCTGAGATCC	CTCTCTCGGG	540
	ATTAGACAGT GG	ACTGTTAT	GGGTGTAGGT	GAATTGGCTT	ATTTTGTCTG	TCCCTGTCTG	600
45	AATGTATTGC AG	GAAYTAAA	AAGGACCAAG	AAGAGGAAGA	AGACCAAGGC	CCACCATGCC	660
	CCAGGCTCAG CA	AGGGAGCTG	CTGGAGGTAG	TAGAGCCTGA	AGTCTTGCAG	GACTCACTGG	720
<b></b>	ATAGATGTTA TT	CAACTCCT	TCCAGTTGTC	TTGAACAGCC	TGACTCCTGC	CAGCCCTATG	780
50	GAAGTICCTT TI	TATGCATTG	GAGGAAAAAC	ATGTTGGCTT	TTCTCTTGAC	GTGGGAGAAA	840
	TTGAAAAGAA GG	GGAAGGGG	AAGAAAAGAA	GGGGAAGAAG	ÄTCAAAGAAG	GAAAGAAGAA	900
55	GGGGAAGAAA AG	AAGGGGAA	GAAGATCAAA	ACCCACCATG	CCCCAGGCTC	AGCAGGGAGC	960
	TGCTGGATGA GA	AAAGRGCCT	GAAGTCTTGC	AGGACTCACT	GGATAGATGT	TATTCAACTC	1020
60	CTTCAGTIGT GI	rtgaactgt	GTGACTCATG	CCAGCCCTAC	AGAAGTGCCT	TTTATGTATT	1080

	GGAGCAACAG CATGTTGGCT TGGCTGTTGA CATGGATGAA ATTGAAAAGT ACCAAGAAGT	1140
	GGAAGAAGAC CAAGACCCAT CATGCCCCAG GCTCAGCAGG GAGCTGCTGG ATGAGAAAGA	1200
5	GCCTGAAGTC TTGCAGGACT CACTGGATAG ATGTTATTCG ACTCCTTCAG GTTATCTTGA	1260
	ACTGCCTGAC TTAGGCCAGC CCTACAGCAG TGCKGTTTAC TCATTGGAGG AMCAKTACCT	1320
10	TGGCTTKKCT CTTGACGTGG ASAAATTGAA AAGAAGGGGA AGGGGAARAA AAGAAGGGGA	1380
10	AGAAGATCAA AGAAGGAAAG AAGAAGGGGA AGAAAAGAAG GGGAAGAA	1440
	CCATGCCCCA GGCTCAGCAG GGAGCTGCTG GATGAGAAAG GGCCTGAAGT CTTGCAGGAC	1500
15	TCACTGGATA GATGTTATTC AACTCCTTCA GGTTGTCTTG AACTGACTGA CTCATGCCAG	1560
	CCCTACAGAA GTGCCTTTTA YRTATTGGAG CAACAGYGTG TTGGCTTGGC TGTTGACATG	1620
20	GATGAAATTG AAAAGTACCA AGAAGTGGAA GAAGACCAAG ACCCATCATG CCCCAGGCTC	1680
20	AGCAGGGAGC TGCTGGATGA GAAAGAGCCT GAAGTCTTGC AGGACTCACT GGATAGATGT	1740
	TATTCGACTC CTTCAGGITA TCTTGAACTG CCTGACTTAG GCCAGCCCTA CAGCAGTGCT	1800
25	GTTTACTCAT TGGAGGAACA GTACCTTGGC TTGGCTCTTG ACGTGGACAG AATTAAAAAG	1860
•	GACCAAGAAG AGGAAGAAGA CCAAGGCCCA CCATGCCCCA GGCTCAGCAG GGAGCTGCTG	1920
30	GAGGTAGTAG AGCCTGAAGT CTTGCAGGAC TCACTGGATA GATGTTATTC AACTCCTTCC	1980
30	AGTTGTCTTG AACAGCCTGA CTCCTGCCAG CCCTATGGAA GTTCCTTTTA TGCATTGGAG	2040
	GAAAAACATG TTGGCTTTTC TCTTGACGTG GGAGAAATTG AAAAGAAGGG GAAGGGGAAG	2100
35	AAAAGAAGGG GAAGAAGATC AAMGAAGRAA AGAAGAAGGG GAAGAAAAGA AGGGGAAGAA	2160
	GATCAAAACC CACCATGCCC CAGGCTCAAC GGCGTGCTGA TGGAAGTGGA AGAGCSTGAA	2220
40	GTCTTACAGG ACTCACTGGA TAGATGTTAT TCGACTCCGT CAATGTACTT TGAACTACCT	2280
	GACTCATTCC AGCACTACAG AAGTGTGTTT TACTCATTTG AGGAACAGCA CATCAGCTTC	2340
	GCCCTTTACG TGGACAATAG GTTTTTTACT TTGACGGTGA CAAGTCTCCA CCTGGTGTTC	2400
45	CAGATGGGAG TCATATTCCC ACAATAAGCA GCCCTTASTA AKCCGAGAGA TGTCATTCCT	2460
	GCAGGCAGGA CCIATAGGCA MGTGAAGATT TGAATGAAAG TACAGTTCCA TTTGGAAGCC	2520
50	CAGACATAGG ATGGGTCAGT GGGCATGGCT CTATTCCTAT TCTCAAACCA TGCCAGTGGC	2580
	AACCTGTGCT CAGTCTGAAG ACAATGGACC CACGTTAGGT GTGACACGTT CACATAACTG	2640
	TGCAGCACAT GCCGGGAGTG ATCAGTCRGA CATTTTAATT TGAACCACGT ATCTCTGGGT	2700
55	AGCTACAAAA TTCCTCAGGG ATTTCATTTT GCAGGCATGT CTCTGAGCTT CTATACCTGC	2760
	TCAAGGTCAK TGTCATCTTT GTGTTTAGCT CATCCAAAGG TGTTACCCTG GTTTCAATGA	2820
60	ACCTAACCTC ATTCTTTGTG TCTTCAGTGT TGGCTTGTTT TAGCTGATCC ATCTGTAACA	2880

	CAGGAGGGAT	CCTTGGCTGA	GGATTGTATT	TCAGAACCAC	CAACTGCTCT	TGACAATTGT	2940
	TAACCCGCTA	GRCTCCTTTG	GTTAGAGAAG	CCACAGTCCT	TCAGCCTCCA	ATTGGTGTCA	3000
5	GTACTTAGGA	AGACCACAGC	TAGATGGACA	AACAGCATTG	GGAGGCCTTA	GCCCTGCTCC	3060
	TCTCRATTCC	ATCCTGTAGA	GAACAGGAGT	CAGGAGCCGC	TGGCAGGAGA	CAGCATGTCA	3120
10	CCCAGGACTC	TGCCGGTGCA	GAATATGAAC	AAYGCCATGT	TCTTGCAGAA	AACGCTTAGC .	3180
10	CTGAGTTTCA	TAGGAGGTAA	TCACCAGACA	ACTGCAGAAT	GTRGARCACT	GAGCAGGACA	3240
	GCTGACCTGT	CTCCTTCACA	TAGTCCATRT	CACCACAAAT	CACACAACAA	AAAGGAGARG	3300
15	AGATATTTTG	GGTTCAAAAA	AAGTAAAAAG	ATAATGTAGC	TGCATTTCTT	TAGTTATTTT	3360
	GARCCCCAAA	TATTTCCTCA	TCTTTTTGTT	GTTGTCATKG	ATGGTGGTGA	CATGGACTTG	.3420
20	TTTATAGAGG	ACAGGTCAGC	TGTCTGGCTC	AGTGATCTAC	ATTCTGAAGT	TGTCTGAAAA	3480
20	TGTCTTCATG	ATTAAATTCA	GCCTAAACGT	TTTGCCGGGA	ACACTGCAGA	GACAATGCTG	3540
	TGAGTTTCCA	ACCTYAGCCC	ATCTGCGGGC	AGAGAAGGTC	TAGTTTGTCC	ATCASCATTA	3600
25	TCATGATATC	AGGACTGGTT	ACTTGGTTAA	GGAGGGGTCT	AGGAGATCTG	TCCCTTTTAG	3660
	AGACACCTTA	CTTATAATGA	AGTATTTGGG	AGGGTGGTTT	TCAAAATTAG	AAATGTCCTG	3720
30	TATTCCRATG	ATCATCCTGT	AAACATTTTA	TCATTTATTA	ATCATCCCTG	CCTGTGTCTA	3780
50	TTATTATATT	CATATCTCTA	CGCTGGAAAC	TTTCTGCCTC	ÄATGTTTACT	GIGCCTITGT	3840
٠	TTTTGCTAGT	GTGTGTTGTT	GAAAAAAAAA	ACATTCTCTG	CCTGAGTTTT	AATTTTTGTC	3900
35	CAAAGTTATT	TTAATCTATA	CAATTAAAAG	CTTTTGCCTA	. ТСААААААА	AAAAAAAAA	3960
	ААААААААА	AAAAAGCGGA	ceceireec				3989

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#### (2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 3735 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

60	GTCGGCGGGC	CSGCTGACGG	GCTTGGCCAG	TCCGCAGCAG	CTGGCTGGGC	CTGCTGTTCG
120	CCCTCTCAAA	GGTAGTGCAN	ATTTTATTCT	GCAGCTGCAG	GAACAGGCAC	GGGTTTGTGT
180	AGAAAACTTG	ATTCCAAAAA	AGAAGTAGTA	CAGGGATTGA	ACTGATGTAA	GGTTGAAGGA
240	CCACAGCTGT	AACAGGGATA	ATCCACAGTA	AGGCACTTGC	GCCGTTCTTC	GGATAAAGTA
300	AATCTCGTTC	TCATCTTTGG	TATGCCAGCA	ATCCTTACCT	TTTCAAGATG	GCCTTATGTG

	ATTTTACTG GCAAAGAAAT CCGGGGAGAA TGTGGCCAAG TITATTATTA ATTCATACCC	360
_	CAAATATTTT CAGAAGGACA TAGCTGAACC TCATATACCG TGTTTAATGC CTGAGTACTT	420
5	TGAACCTCAG ATCAAAGACA TAAGTGAAGC CGCCCTGAAG GAACGAATTG AGCTCAGAAA	480
	AGTCAAAGCC TCTGTGGACA TGTTTGATCA GCTTTTGCAA GCAGGAACCA CTGTGTCTCT	540
10	TGAAACAACA AATAGTCTCT TGGATTTWTT GTGTTACTAT GGTGACCAGG AGCCCTCAAC	600
	TGATTACCAT TTTCAACAAA CTGGACAGTC AGAAGCATTG GAAGAGGAAA ATGATGAGAC	660
1.5	ATCTAGGAGG AAAGCTGGTC ATCAGTTTGG AGTTACATGG CGAGCAAAAA ACAACGCTGA	720
15	GAGAATCTTT TCTCTAATGC CAGAGAAAAA TGAACATTCC TATTGCACAA TGATCCGAGG	780
	AATGGTGAAG CACCGAGCTT ATGAGCAGGC ATTAAACTTG TACACTGAGT TACTAAACAA	840
20	CAGACTCCAT GCTGATGTAT ACACATTTAA TGCATTGATT GAAGCAACAG TATGTGCGAT	900
	AAATGAGAAA TTTGAGGAAA AATGGAGTAA AATACTGGAG CTGCTAAGAC ACATGGTTGC	960
25	ACAGAAGGTG AAACCAAATC TTCAGACTTT TAATACCATT CTGAAATGTC TCCGAAGATT	1020
25	TCATGTGTTT GCAAGATCGC CAGCCTTACA GGTTTTACGT GAAATGAAAG CCATTGGAAT	1080
	AGAACCCTCG CTTGCAACAT ATCACCATAT TATTCGCCTG TTTGATCAAC CTGGAGACCC	1140
30	TTTAAAGAGA TCATCCTTCA TCATTTATGA TATAATGAAT GAATTAATGG GAAAGAGATT	1200
	TTCTCCAAAG GACCCGGATG ATGATAAGTT TTTTCAGTCA GCCATGAGCA TATGCTCATC	1260
35	TCTCAGAGAT CTAGAACTTG CCTACCAAGT ACATGGCCTT TTAAAAAACCG GAGACAACTG	1320
55	GAAATTCATT GGACCTGATC AACATCGTAA TTTCTATTAT TCCAAGTTCT TCGATTTGAT	1380
	TTGTCTAATG GAACAAATTG ATGTTACCTT GAAGTGGTAT GAGGACCTGA TACCTTCAGC	1440
40	CTACTTTCCC CACTCCCAAA CAATGATACA TCTTCTCCAA GCATTGGATG TGGCCAATCG	1500
	GCTAGAAGTG ATTCCTAAAA TTTGGAAAGA TAGTAAAGAA TATGGTCATA CTTTCCGCAG	1560
45	TGACCTGAGA GAAGAGATCC TGATGCTCAT GGCAAGGGAC AAGCACCCAC CAGAGCTTCA	1620
73	GGTGGCATTT GCTGACTGTG CTGCTGATAT CAAATCTGCG TATGAAAGCC AACCCATCAG	1680
	ACAGACTGCT CAGGATTGGC CAGCCACCTC TCTCAACTGT ATAGCTATCC TCTTTTTAAG	1740
<b>50</b> .	GGCTGGGAGA ACTCAGGAAG CCTGGAAAAT GTTGGGGCTT TTCAGGAAGC ATAATAAGAT	1800
	TCCTAGAÁGT GAGTTGCTGA ATGAGCTTAT GGACAGTGCA AAAGTGTCTA ACAGCCCTTC	1860
55	CCAGGCCATT GAAGTAGTAG AGCTGGCAAG TGCCTTCAGC TTACCTATTT GTGAGGGCCT	1920
J.	CACCCAGAGA GTAATGAGTG ATTTTGCAAT CAACCAGGAA CAAAAGGAAG CCCTAAGTAA	1980
	TCTAACTGCA TTGACCAGTG ACAGTGATAC TGACAGCAGC AGTGACAGCG ACAGTGACAC	2040
60	CAGTGAAGGC AAATGAAAGT GGAGATTCAG GAGCAGCAAT GGTCTCACCA TAGCTGCTGG	2100

	AATCACACCT (	GAGAACTGAG	ATATACCAAT	ATTTAACATT	GTTACAAAGA	AGAAAAGATA	2160
5	CAGATTTGGT (	GAATTTGTTA	CTGTGAGGTA	CAGTCAGTAC	ACAGCTGACT	TATGTAGATT	2220
5	TAAGCTGCTA	ATATGCTACT	TAACCATCTA	TTAATGCACC	ATTAAAGGCT	TAGCATTTAA	2280
	GTAGCAACAT '	TGCGGTTTTC	AGACACATGG	TGAGGTCCAT	GGCTCTTGTC	ATCAGGATAA	2340
0	GCCTGCACAC	CTAGAGTGTC	GGTGAGCTGA	CCTCACGATG	CTGTCCTCGT	GCGATTGCCC	2400
	TCTCCTGCTG	CTGGACTTCT	CCCTTTGTTG	GCCTGATGTG	CTGCTGTGAT	GCTGGTCCTT	2460
15	CATCTTAGGT	GTTCATGCAG	TTCTAACACA	CTTCCCCTTC	GGTCAATAGT	TTCCCAATTT	2520
()	CAGGATATTT	CGATGTCAGA	AATAACGCAT	CTTAGGAATG	ACTAAACAAG	ATAATGGCAG	2580
	TTTAGGCTGC	ACAACTGGTA	AAATGACTGT	AGATAAATGT	TGTAATTAGT	GTACACGTTT	2640
20	GTATTTTTGT	TAATATAGCC	GCTGCCATAG	TTTTCTAACT	TGAACAGCCA	TGAATGTTTC	2700
	ATGTCTCCCT	TTTTTTTTG	TCTATAGCTG	TTACCTATTT	TAGTGGTTGA	AATGAGAGCT	2760
25	AGTGATGACA	GAAGGATGTG	GAATGTCTTC	TTGACATCAT	TGTGTATTGC	TGGTAATCAA	2820
23	GTTGGTAACG	ACTACTTCTA	GCAGCTCTTA	CCACTATGAC	TTAAGTGGTC	CTGGAAGGCA	. 2880
	GTAAGTGGAG	GTTTGCAGCA	TTCCTGCCTI	CATGAGGGCT	TCTACCACTG	ACCACTTTGC	294Ò
30	ACGTACCTGG	CTCCCAGATT	TACTTAGGTA	CCCCACGAGT	CGTCCACATA	AGCAGCTTCA	3000
	TCTTTACCTT	GCCAGAGTTG	ACAATTATGG	GATACTCTAG	TCTACTTATA	CTTGTGTTCC	3060
35	CATCTGTCTG	CCATCCTCTG	AAGGCCAGGA	CCCAGȚCATA	CATCCTTAGA	AACCAAAGTA	3120
<i>33</i>	TGGTTTTTGT	TTTCTCTTGG	AATGTCAGGT	CTTAAGGCAT	TTAATTGAGG	GACAAAAAA	3180
	AAAAAAAGCC	GATATAGTAG	CTAGCTACTT	AAGCATCCAT	GGTATTGCT	CCATATCAAA	3240
40	GCAGATTTGC	AGGACAGAAA	GAGTAAATT	A GCCTTCAGTC	TTGGTTTACA	GCTTCCAAGG	3300
	AGAGCCTTGG	CCACCTGAAA	TGTTAACTCC	GTCCCTTCCI	GICTCTAGTI	CATCAGCACC	3360
45	TGCAGATGCC	TGACTCTTGT	TAGCCTTACT	T ATTCAATACA	GTCCTTAGAT	TCACGGTATG	3420
73	CCTCTTCCTA	TCCAGGCACC	TATTCTGAA:	CACCATGITC	CTCTGCAGCT	AGAGTTGATA	3480
	GGAGAAAATC	CATTTGGGTA	GATGGCCTA	r gaatttgtac	TAGACTTTC	AAATGAGTGA	3540
50	TTTGTTAGCT	TGGTACTTTT	AAGTTIGIG	G ȚACAGATCC	CCAAACCCA	r ACTCTGAGCA	3600
	ATTAACTGCC	TTGAACATAG	AGAAAATTA	A GGCCTCACAC	GATGAGTCT	CATTCTCTGT	3660
55	AAATGCTTAT	TTTATCATAG	TCTTTAGCC	N CTACTATGA	TAAAAT	TCTTCNGCCG	3720
<i>,,</i>	GGTGTGGTGA	CTCAC				•	3735

#### (2) INFORMATION FOR SEQ ID NO: 30:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1667 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

10 TAGTAATTCA TITAACTCCT CTTACATGAG TAGCGACAAT GAGTCAGATA TCGAAGATGA 60 AGACTTAAAG TTAGAGCTGC GACGACTACG AGATAAACAT CTCAAAGAGA TTCAGGACCT 120 GCAGAGTCGC CAGAAGCATG AAATTGAATC TTTGTATACC AAACTGGGCA AGGTGCCCCC 180 15 TGCTGTTATT ATTCCCCCAG CTGCTCCCCT TTCAGGGAGA AGACGACGAC CCACTAAAAG 240 CAAAGGCAGC AAATCTAGTC GAAGCAGTTC CTTGGGGAAT AAAAGCCCCC AGCTTTCAGG 300 20 TAACCTGTCT GGTCAGAGTG CAGCTTCAGT CTTGCACCCC CAGCAGACCC TCCACCCTCC 360 TGGCAACATC CCAGAGTCCG GGCAGAATCA GCTGTTACAG CCCCTTAAGC CATCTCCCTC 420 CAGTGACAAC CTCTATTCAG CCTTCACCAG TGATGGTGCC ATTTCAGTAC CAAGCCTTTC 480 25 TGCTCCAGGT CAAGGAACCA GCAGCACAAA CACTGTTGGG GCAACAGTGA ACAGCCAAGC 540 CGCCCAAGCT CAGCCTCCTG CCATGACGTC CAGCAGGAAG GGCACATTCA CAGATGACTT 600 30 GCACAAGTTG GTAGACAATT GGGCCCGAGA TGCCATGAAT CTCTCAGGCA GGAGAGGAAG 660 CAAAGGGCAC ATGAATTATG AGGGCCCTGG AATGGCAAGG AAGTTCTCTG CACCTGGGCA 720 ACTGTGCATC TCCATGACCT CGAACCTGGG TGGCTCTGCC CCCATCTCTG CAGCATCAGC 780 35 TACCTCTCTA GGTCACTTCA CCAAGTCTAT GTGCCCCCCA CAGCAGTATG GCTTTCCAGC 840 TACCCCATTT GGCGCTCAAT GGAGTGGGAC GGGTGGCCCA GCACCACAGC CACTTGGCCA 900 40 GTTCCAACCT GTGGGAACTG CCTCCTTGCA GAATTTCAAC ATCAGCAATT TGCAGAAATC 960 CATCAGCAAC CCCCCAGGCT CCAACCTGCG GACCACTTAG ACCTAGAGAC ATTAACTGAA 1020 TAGATCTGGG GGCAGGAGAT GGAATGCTGA GGGGGTGGGT GGGGGTGGGA AGTAGCCTAT 1080 45 ATACTAACTA CTAGTGCTGC ATTTAACTGG TTATTTCTTG CCAGAGGGGA ATGTTTTTAA 1140 TACTGCATTG AGCCCTCAGA ATGGAGAGTC TCCCCCGCTC CAGTTATTGG AATGGGAGAG 1200 50 GAAGGAAAGA ACAGCTTTTT TGTCAAGGGG CAGCTTCAGA CCATGCTTTC CTGTTTATCT 1260 ATACTCAGTA ATGAGGATGA GGGCTAGGAA AGTCTTGTTC ATAAGGAAGC TGGAGAACTC 1320 AATGTAAAAT CAAACCCATC TGTAATTTCG AGTGGGTGGA GCTCTTGCTT TTGGTACATG 1380 55 CCCTGAATCC CTCACTCCCT CAAGAATCCG AACCACAGGA CAAAAACCAC CTACTGGGCT 1440 CTCTCCTACC CTGCCCTCCT CCCTTTTTT TACCCCTCTC TTTTTTATTT TTTCTTTGCT 1500 60

	CTTTAGAACC	CAGTGAAAAA	TACCAGGGTA	CTGGGGTGCA	ACTCTTTCTT	ATGATAGGTC	1560
	ATTAGTGCTT	TAAGCAAAAG	ATATTAGCAG	CTTTGACTGC	AGCATTAGCA	ATTAGGRAAA	1620
5	AAAAAANWA	AAAACTCGAG	GGGGGGCCCG	GTTACCCAAT	TCGCCCT		1667

# 10 (2) INFORMATION FOR SEQ ID NO: 31:

# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1408 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:

	1	
20	ATTACACACC TGAGCACTGT GCCTGGCAAG ACCTGTCTTA ATAGATTAGA GAACCACTGA	60
	TAGATGGTCA GCTTTCTGTA GCAGTGAGAA CCCTACATTT CAAATGTGGA TAGCACCTTT	120
25	GCGGGGAAAC ATCACTTGGC ACATCTGCAT TCTTTTTTGA CACAGGGTCT CACTCTGTTG	180
23	CCCAGGCTAG AGTGCATGGC ACGATCTTAG CTCACTGCAA CCTCCACCTC CCAAGTTCAA	240
	GCGATTCTTC TGCCTCAGCC TCCTGAGCAG CTGGGATCAC AGACATGCGC TACCATGCCC	300
30	AGCTAATTTT TTGTATTTTT TGTKTGTTTG TTTTTGTTTK TAAGTAGAGA CGGGCTTTCA	360
	CCACGITGGS CAGGCAGGTC TCGAACTCCT GAMCTCAGGT GATCCACCCA CATCTGCGTT	420
35	CCAATATCTT TCTCAACATA ATGATAGCCG TAATTAATAT TTTCCAGTAC ATTTTTATGC	480
33	CTTTACACAC GAGAGTGGTA GACAGACACA AACCCAGATC TGTCTGACTC CAAAGCCCGT	540
	TTGTCATCAT TCCTTTTACG GTATCCTATA GTGGTATCCT TTACAGAAAG ACAGCTTTTA	600
40	CCCAACAAAG ACTTAACTTC CCAGGATGCC AGAAGGACAA AGCGGGATTG CTTTTAAGRA	660
	GRAAGTTATC AAGAMCTTAT TITATAAATG AGATTAGATA GGGAAAGGCA ATTTATCTTT	720
45	ATTAAAAACT GAAAAGGCCA GCATAGGGAA GGAGGTCCTT CGGTGGTCTT TTTCAGGGAA	780
43	ATACTTCAGT TGCTTTTATT AGAAACAGAT AGTACCTAAG GTTTTGAGGT AGGWACAGCT	840
	TAAGGCATGC TAATGKTCAT GGGTCCTTCC ATAGTCATTT TKGTATTTTG GTTWACATTT	900
50	GAGCAATAGG CAGCCCTTCA CTGCTGCTGG AYTCATTCCT GCCAYTATTA CAGGTGACAG	960
	AGGAGACAGG AGGTATGTCT TTTCTATTTT TAWACATGCT TTATATTTAA CACAAGCTCT	1020
55	TGGGTATCTT AGATAAACAG AAGTTGCCTA GCACTCCTTT TAGTGCATTG AACCCTTTAA	1080
55	CATTTAAGCA AAATAATAAA CAGTCTTTTG AGGTTCCTTA ACAATGAAAC GTGTTCGAGT	1140
	GGCAGCAGCG GAATCCATGC YTCTTCTCCT GGAGTGTGCA AKAGTCCGTG GTCCTGAGTA	1200
60	TCTCACACAG ATGTGGCATT TTATGTGTGA TGCTCTAATT AAGGCCATTG GTACAGAACC	1260

AGATTCAGAC	GTCCTCTCAG	ARATAATGCA	TTCTTTTGCA	AAGGTGAATA	TTTTTCTCTT	1320
AAAAAATATG	TATAAGGTGG	TATGTTCATT	TATTAGTCTT	GCTAAAAAA	AAAAAAAA	1380
ACTINGAGGG	GGGGNCCGGT	ACCCAATT				1408

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# (2) INFORMATION FOR SEQ ID NO: 32:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGIH: 2031 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLCGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 32:

	(XI) SEQUENCE DESCRIPTION. BEG ID NOT DE-	
20	AGGATATGCA TGATTCTTAA CCAGGCTATA TGTTAAAAAA AAATTGGAAA ATGCAATACA	60
٠	TTTTTTATTA TACAAACTAC AGAATGAGTA TGCAAGTTTT ATTTATCAAA ATGTAATGGA	120
25	TTTTTAAAGG CTGAGAAATT TTCCTTATAC CTACCTTTTC AGTTATTTTA ATTATACCAA	180
	ATTATCAACT AGAATAGCTT CATCCATATG AAATATAAAA TGAAGAGACA CCTAGGCTCT	240
30	ATCAGGCTTA GGATTCTTTG AACTTATTTC CACTTTAATT TCTCAGTGGA AGTTAAGAGG	300
30	GGTGAGAAAA CAAAGAAGGG GAAAAACTGA CAACTAACAA AACCAGCACC ACATCGCTAG	360
	GTGGTGCTTA CTAATTACCT TCTCAGGATT TTCCTCAGAT TGAAAAGCTT ATGAGGATTT	420
35	CTTGGGAGTC TTAATAACCT GCCTGTTAGT ACAGAGCTTT CCTGATGATA TTTACTCTTG	480
٠	AGCACATGTG GTTGTAAAAC CTTAACTTTC TTTCTCCAGG AGGGTGGTGA TAGAAACAGA	540
40	TOGTAGTATT TATGAACTGA TGTTCTCGTG AAATGTTGAG GGTGGGGAGA AAAGACTTTA	600
40	AGGGAGGAGA GCCATCTATT TTGTTCCTAA AGCCACCTCT CAGCAGAATC GTCATGTTTT	660
	TCTGATGCAC CGCTCTGCTT CATGCCCAAG ATGACTTGCG AGGCAATCTC AGGAGCTGTG	720
45	GACTTAACCR TTGCAAAGCA CACTGTCTTT CTCAGCGTTC TCTGCAAGTC AGTAGGTGTT	780
	AGTATGGTTG CAAAGTTCAC TGTCTCAGCA AAGTTGAACT GGGCTACCTC TCTACAGCTG	840
50	TTTCCTCAGA GGGAAAAATC TTGAGACCAG ATGGTGGAGC TCTGGAGTCA GAGGAAATGG	900
50	GTGTCTTCAG CACAAAGCTG CTGCTTTTAC TTCAGCCACT TCTGACATTT TTACATACCG	960
	AGCCTGAGAT TRIGIGATTA TCTCAAATCA AATCACTTIG ATGGAGATAA ATAATCAAAA	1020
55	CTGTTTEATA GTCATTGATT TGGTGAGAAC AGTAATGGAA AATGGTGTTG AAGGACTTCT	1080
	CATTTTTGGA GCTTTCCTTC CAGAGTCCTG GCTGATTGGT GTTCGCTGTT CATCTGAGCC	1140
60	CCCAAAAGCA TTATTACTGA TACTTGCACA CAGTCAAAAG CGCAGACTGG ATGGATGGTC	1200

•	TTTTATAAGG	CATTTAAGGG	TACACTACTG	TGTTTCACTG	ACCATACATT	TTTCTTAGCC	1260
	CCTCAAGTAA	TATAGCACAG	AGTTATGAAT	GACAATTCCC	CTAACCATTC	CTCTTCATAT	1320
5	CTGCCTCTTC	CCCTTACCAT	CGTAATTCTC	CAAACTGGTC	ATAAAGGCAC	TCTGTGAAGA	1380
	TATTGGGGAC	TGACATCTTA	AGCTCTCACC	TGGCTGCAGT	AGGAAAGGCC	AAACTGACGA	1440
10	СААААААА	ATTCTTTATA	AAGATGATAT	GGTAACATGT	ATCTTTGCCC	TGGGTCTGGG .	1500
,	TGGGTCCAGT	CAGTCTCAGA	TTTACAAGCA	TTTAGGAGCC	TAGGTAAAAG	CTGCTAGTAT	1560
	TCTTTTAAAA	GTTACATTTA	TGACTTGCAA	TGATAGAAAA	CTCCTTCCAA	TTAAATGGCA	1620
15	TTTTATAATA	TTATGTGTGT	ACTICACAGT	GTTAAAAATA	CCCTCATACG	TTATTGCATT	1680
	TGATCTTCAC	AGAAAGTGCA	TTTTAACCAG	TACTCTGGGT	GCAATAAATA	ATATGTAGAA	1740
20	ATTTAAGTCC	TCCAATTCCA	GCATATCCAG	TGAGTTTTGA	CAGTGTGTTT	ATGTGGAATG	1800
20	TTTAAGGATA	TACAATTGTA	CTTTATATAA	ATTGGTTCTT	GTTCTTCTTA	AATGTGACAT	1860
	GAAATAATTG	TGCTGCTACA	TTATACTGGA	AATTAACAGG	GGAAAAGGGA	AGAGCTCTTG	192
25	GCTCCCTTGA	GGTTCTGCTA	GTGGTGTTAG	GAGTGGTTAC	AACTGAGCTT	TTAGTAACCA	198
	TTTAACCGTA	TGTAAACTTG	GTTTCTAATI	TAAAAAAAA	TTCTTTTTCC	. A	203

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## (2) INFORMATION FOR SEQ ID NO: 33:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 971 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

					· -	•	
. 60	GAGGGAGAGC	TGACACGCGG	GGGCGCGAG	GGACATCCAC	CTCGGCCGCG	CGCGTCGGAA	
120	CATTGTTTTC	TATTCAGATT	ATGCATTTCT	GCCAAAAACC	TGGAGCCGAT	AGTGTTCTGC	
180	CACCGAAGAA	AAGAGGAGAG	AGACAAAAGA	TACTGCTCAG	GGGGCCTTTT	TTTTATCTGT	
240	GAAGGGAGAC	AGACAAGCAA	AACTGCTCTA	TCGTCCAGAA	AAGTTTTGCA	GTGAAAATAG	
300	CTACTGCAGC	GCTCGAAATT	GCTAAAGACG	COGCTACCTG	CCCATTATGA	CTACTAAATG	
360	AGTCATAAAA	GTGTTGGGCA	TTTGTTCTTG	CCCCAAATGG	ATGAAGGCCA	CGGACACAAA	
420	AGTTATACCC	. AGCGAAAAGT	CCTGGAGAAA	AGATATGTGC	TTGCTATGAC	GGCCTAGACA	
480	GGATGCTACA	AGATTCCACC	GCAGAAGGCA	GGAAGGCTAT	CATACGGAAA	CCTTCATTTG	
540	TGAGACATTT	CACGGAGCAT	ACCAAAGGAC	TTATGCTGTG	G AGATTGAACT	TTGATTTTTG	
600	CCTCTACTTG	CCGAGATAAA	CTCTCTAAAG	A TGACAGGCAG	ACATGGACAA	AAACAAATAG	

•	CAAAGGGAAT TTGAAAAAGA TGAGAAGCCA CGTGACAAGT CATATCAGGA TGCAGTTTTA	660
<b>.</b>	GAAGATATTT TTAAGAAGAA TGACCATGAT GGTGATGGCT TCATTTCTCC CAAGGAATAC	720
5	AATGTATACC AACACGATGA ACTATAGCAT ATTTGTATTT CTACTTTTTT TTTTTAGCTA	780
	TTTACTGTAC TTTATGTATA AAACAAAGTC ACTTTTCTCC AAGTTGTATT TGCTATTTTT	840
10	CCCCTATGAG AAGATATTTT GATCTCCCCA ATACATTGAT TTTGGTATAA TAAATGTGAG	900
	GCTGTTTTGC AAACTTAAAA AAAAAWWAAA AAAACTSGAG GGGGGCCCGT ACCCAANTCG	960
15	CCGNATATGA T	971
13	(	
20	(2) INFORMATION FOR SEQ ID NO: 34:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1792 base pairs	
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:	•
30	GAACCCCCTT TCTCCTGGTA AAGGGTAAGG GGGGGGATAA TGTTTACCAC AGGTACGAAA	. 60
J.O	TAGTCACTTT AACATTGAGA CCTCTGCCTC ATTGAATTCA GGTTTTTTAA GTACTTGAAA	120
	CTCTTCAGAT TCTCCTTATT TTAGTTTCTT TTTACATTTA TGAAGTAGAA AGCATTGTTT	180
35	TGTAAACTGT TTTGAAAATA AATAGCCTAG TCTCTTATCC TCTTTAGCGT GGATTAAAGG	240
	TGAAGTTCTG CAAATGGGAG AGTGTTCACA GTAGATAGCT CAGATTGATT GAACACATTT	300
40	GAGGAAGAGA CTCCTGCATG AGATACCAGC ATTTTTACAA ATACTTTTTA TGTACATTCT	360
	TTATTTTGTC ATTTTGTCAA CCCTCTCCCC AAGCACATCT TCTTTCCTTT TACTATGTCT	420
	ATGTAGGGAA AAACAAAACA AAAAATTGCA CTTACGTTAC ACTCCCAAAA TGTGGGTAAT	480
45	CCGTGTCTTT CAAAAAACAT TTCTGTTTTT TGTTTTGTTT	540
	TGACAAGTTT GGGTGCTTGT GGCACGTATG TATGAAGCGG GAGGGGGATG ASAATTGCCT	
50	GTCCTTCAGT ARGCTGTAAA AGTAATTTAC ATGTAAGTAA AAAGGGAAAA TAGAATAGAT	660
	GCCAAAGTCA TTTATTCAGT CCTTAGTTTT CTTATGTGGC ATTACTGCAT CTGCTAGTTA	720
	GTGAGAAAGC ACCCTCAGCT TTTACTGCTC CCCTCCCTGC CTGCCAACAC ACTTGATGTG	780
55	TGCAAACAGC CCTCAAGTAT CTGTCAGATG ACCTATATAA GGTATTGAAT AAGGTATTCT	٠.
	TGTCAGTTTA GAAATGGACT GGATAAAACT TACTTGGTTG TCATTATTTT ATCTCATTTG	900
	TCCTGTTACA TGCCCTATGT TAAGATAATT ATATTGCCAC TAATAATCAA GATGCTAAAT	960

•	GAGTATTACA	ACTGGCTAAT	ATCATTTTT	ATATACAAGG	GTATGTGTAT	ATTTGGAATT	1020
	GRTATGAGAA	ACTCATTIGT	ACCCATTTGA	GTGATATTGC	ACAACAAACA	CAGATAÝCTA	1080
5	CAGACTCCGT	TTTCATTTTC	TCGTGTTCTT	TATGATAATG	ATCTTTGTAG	ATTGGTTATT	1140
	TCTGTACTTT	ATCTGTAATA	AACTTTGTAG	ATCCTGTGAA	CCATTACTTT	GCCTAAATCA	1200
0	CTTGAGACTT	GAGTCTTTAA	TAACAAAGCA	TCAATATTCA	CTAAAGTCAA	TCTCTTTTGA	- 1260
.0	GTTTCTGTGA	CTTGGCTAGA	AGCTCTTGAC	ACTAAGGGAT	TAGTGTTAAT	TTTCCCTGGG	1320
	GGTGTTCCAC	TAGGGCATTA	CTGTATAATG	ACTTGATGTT	GCCACATAGA	CTTCAAGATA	1380
5	TATAATATTT	TGAGGATTTT	GTTGATTGGC	CTATGTTTTA	TTGCATAGTG	TGAAACGTGT	1440
	AAAGCTTGGT	TAACCTGTAT	ATAGATAGCT	TATTGTTGAC	TAGTTATAGT	GTATTTAGGG	1500
20	TTGCCTGTAA	TATTTAAGCT	TCTTTACTGA	TGTGTGTGCT	GGTAGGAACA	TATAATTTTT	1560
20	GTACATTATA	TTTACTGAGA	TGTTGCCTTT	TTTATTTTAC	AAATACTTTG	GAATTCCAAT	1620
	GTGTTTTTG	CTTCCGTGAG	GATTAATTIG	GAAAGGTTTT	TAATGACATT	CCACTGATTT	1680
25	CAGATTTTGC	TTGAGATTGA	CTTCAATAAA	TTGTCCTGTA	TGTTCCAAAA	AAAAATTAAA	1740
	AAACTCGAGG	GGGGCCCGGT	ACCCAANNCG	CCGGATATGA	TCGTAAACAA	TC	1792

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# (2) INFORMATION FOR SEQ ID NO: 35:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 896 base pairs

- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

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#### (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:

	AGTTGNANAC	AACAGGACCT	GAGTCCTTGG	GCAGCACCAG	TAGGTTGCCC	CYTGCYTCYT	60
45	GCCAGCYTCA	CYTGCCACYT	TYTGCCCCTY	TCGGGATGCC	TTCGCAGACA	GAGYTYTTCG	120
73	CTGCCTGTGG	TGGCCAYTCT	TIGCTTTIGG	TTYTCTTGCC	CCTTGGCCTC	CCTTTTTGTC	180
	CCCGGGCAGC	CTTGTGTGAC	CTGCCCTTTT	CCCTCCCTTC	CTTTCCAGGA	CAAGCACGCC	240
50	GAGGAGGTGC	GGAAAAACAA	GGAGCTGAAG	GAAGAGGCCT	CCAGGTAAAG	CCTAGAGGCC	300
	AAAGAACTTT	CCAGGTCAGC	CGGACAGCTC	CAGCAGCTCC	ACGTTCCAGG	CAGCCTCGMC	360
55	CGCCGGCTGC	GCTCCCAGCA	CIGGGGITTG	GGGGGAGGGG	GGTGGCCAAG	GGGCGTTTCC	420
<b>33</b>	TCTGCTTTTG	GTGTTTGTAC	ATGTTAAGAA	TTGACCAGTG	AAGCCATCCT	ATTTGTTTCC/	480
	GGGGAACAAT	GACGGGGTGG	GARAGGGGAG	AGGAGAGAGT	TTGGGAAAGG	GAGATGGAGA	540
60	AGAACTCAAG	GACATTGCAA	CCCTGCCCGG	CGCAGATCTG	ATTTTCACAT	CTCTACCTGG	600

•	ACATTGAGCC TCCCAGGCAC CATGTTGAGG AGAGATGAAA ACCAGGGCGG TAGAACTTCA	660
5	GGGTGAAGGA CAGAGTCCTG GGTGGGGCAG CGGCTGCAGG GCGCACCAGA GAACCCAGCC	720
3	AGAGGGGGTG TGAGTACCAG TGGTGTTGCT TCCACCCTGC AGCAGGTGGG ATGAGGTCTG	780
	TGTGTGTGT TGAACCATCA TTTTTTGATC ATCATGACCA ATGAAACATT GAAAAAAAAA	840
10	AAAAAAACTG GAGGGGGCC CGTACCCAAN TCGCCGNATA GTGATCGTAA ACAATC	896
	,	
15	(2) INFORMATION FOR SEQ ID NO: 36:	•
20	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 912 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:	
25	TCGACCCACG CGTCCGGTCA GCCAGTCGCA TCCAGCCATG ACAGCCTTCT GCTCCCTGCT	60
•	CCTGCAAGCG CAGAGCCTCC TACCCAGGAC CATGGCAGCC CCCCAGGACA GCCTCAGACC	120
30	AGGGGAGGAA GACGAAGGGA TGCAGCTGCT ACAGACAAAG GACTCCATGG CCAAGGGAGC	180
50	TAGGCCCGGG GCCAKCCGCG GCAGGGCTCG CTGGGGTCTG GCCTACACGC TGCTGCACAA	240
	CCCAACCCTG CAGGTCTTCC GCAAGACGGC CCTGTTGGGT GCCAATGGTG CCCAGCCCTG	300
35	ARGCAGGGA AKGTCAACCC ACCTGCCCAT CTGTGCTGAG GCATGTTCCT GCCTACCATC	360
	CTCCTCCCTC CCCGGCTCTC CTCCCAGCAT CACACCAGCC ATGCAGCCAG CAGGTCCTCC	420
40	GGATCACYGT GGTTKGGTGG AGGTCTGTCT GCACTGGGAG CCTCARGARG GCTCTGCTCC	480
40	ACCCACTTGG CTATGGGAGA GCCAGCAGGG GTTCTGGAGA AAAAAACTGG TGGGTTAGGG	540
	CCTTGGTCCA GGAGCCAGTT GAGCCAGGGC AGCCACATCC AGGCGTCTCC CTACCCTGGC	600
45	TCTGCCATCA GCCTTGAAGG GCCTCGATGA AGCCTTCTCT GGAACCACTC CAGCCCAGCT	660
	CCACCTCAGC CTTGGCCTTC ACGCTGTGGA AGCAGCCAAG GCACTTCCTC ACCCCYTCAG	720
50	CGCCACGGAC CTYTYTGGGG AGTGGCCGGA AAGCTCCCSG GCCTYTGGCC TGCAGGGCAG	780
50	CCCAAGTCAT GACTCAGACC AGGTCCCACA CTGAGCTGCC CACACTCGAG AGCCAGATAT	840
	TTTTGTAGTT TTTATKCCTT TGGCTATTAT GAAAGAGGTT AGTGTGTTCC CTGCAATAAA	900

CTTGTTCCTG AG

	(1)	SECUENCE	CHARACTERISTICS
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(A) LENGTH: 1382 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 37:

	(X1) SEQUENCE DESCRIPTION. SEQ 25 No.	
10	AATTCGGCAC GAGCGGAGGC GAGCGAAACT RAGGGCGAAA GTTGTGTGTC GTGTTGGCAG	60
	GAGGGCCTAG AAGGGAAAGA CTGTCTAGTG GGACAATGTC ATATTATAAA TTTGGAATGC	120
15	TGAATAGAAA ATTATAGATT TIGATATTGA AGGAAATGAA GCGAAGCYTA AATGAAAATT	180
15	CASCTCGAAG TACAGCAGGC TGTTTGCCTG TTCCGTTGTT CAATCAGAAA AAGAGGAACA	240
	GACAGCCATT AACTTCTAAT CCACTTAAAG ATGATTCAGG TATCAGTACC CCTTCTGACA	300
20	ATTATGATTT TCCTCCTCTA CCTACAGATT GGGCCTGGGA AGCTGTGAAT CCAGAGTTKG	360
	CTCCTGTAAT GAAAACAGTG GACACCGGGC AAATACCACA TTCAGTTTCT CGTCCTCTGA	420
25	GAAGTCAAGA TTCTGTCTTT AACTCTATTC AATCAAATAC TGGAAGAAGC CAGGGTGGTT	480
23	GGAGCTACAG AGATGGTAAC AAAAATACCA GCTTGAAAAC TTGGRATAAA AATGATTTTA	540
	AGCCTCAATG TAAACGAACA AACTTAGTGG CAAATGATGG AAAAAATTCT TGTCCAATGA	600
30	GTTCGGGAGC TCAACAACAA AAACAATTAA GAACACCTGA ACCTCCTAAC TTATCTCGCA	660
	ACAAAGAAAC CGAGCTACTC AGACAAACAC ATTCATCAAA AATATCTGGC TGCACAATGA	720
35	GAGGGCTAGA CAAAAACAGT GCACTACAGA CACTTAAGCC CAATTTTCAA CAAAATCAAT	780
33	ATAAGANACA AATGTTGGAT GATATTCCAG AAGACAACAC CCTGAAGGAA ACCTCATTGT	840
	ATCAGTTACA GTTTAAGGAA AAAGCTAGTT CTTTAAGAAT TATTTCTGCA GTTATTGAAA	900
40	GCATGAAGTA TIGGCGIGAA CATGCACAGA AAACTGTACT TCTTTTTGAA GTATTAGCTG	960
	TTCTTGATTC AGCTGTTACA CCTGGCCCAT ATTATTCGAA GACTTTTCTT ATGAGGGATG	1020
45	GGAAAAATAC TCTGCCTTGT GTCTTTTATG AAATCGATCG TGAACTTCCG AGACTGATTA	1080
40	GAGGCCGAGT TCATAGATGT GTTGGCAACT ATGACCAGAA AAAGAACATT TTCCAATGTG	1140
	TITCTGTCAG ACCGGCGTCT GTTTCTGAGC AAAAAACTTT CCAGGCATTT GTCAAAATTG	1200
50	CAGATGTTGA GATGCAGTAT TATATTAATG TGATGAATGA AACTTAAGTA GTGATAAAAG	1260
	GAAGTTTAGC ATAAATTATA GCAGTTTTCT GTTATTGCTT AATTTACCAT CTCCATAGTT	1320
55	TTATAGCTAC TATTGTATTT CACTTGTTGA ATTAAAGTAT TTGAATTCTT TTAAAAAAAA	1380
JJ	AA	1382

	(2) INFORMATION FOR SEQ ID NO: 38:	
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 872 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38:	-
	GGGCTACTTC AAAGCCCTGG GCCTTATTTC TTCAGGTAAA AAAATATAAA GTCAGATCTC	60
	ATCCCGGCTG GCCATGCTGT TAGACCCTTT CATCCTTCTC TTCTGCCTCT TCTCAACAGC	120
15	TGCCCAGTCC TGTTTGGAAT TCATATACAT ACAGTTCTAA TACTGATGTA TTTACCCTCA	180
	TAAGCCACTC AACCCAGAAT CTTATTTGAA TTATAATCCA GAAACATCAG GTGACGTGTG	240
20	AGACTACTGT ATGAGAAAGA GACAGTTTAA GGGTCAGTCC AATGGAAAAA AGAGTTCTCA	300
20	GAGCTTTCTT TAGCTTATTC TCATCAAAGA GCTTTCTCTG CAGAAGGAAC CTACTGGTTC	360
	CTCCTTTCCA GTCCTAGAAA TCCTGACCTA GAGTGGCTTA ATCCTGCTAG CACCTCTCTC	420
25	TCGCACTCTG GTGCCAAATG ACTCCAGGAA CTGGGCCCATG ATGTGGTGGG AATGACCTTA	480
	CCCTGAGCAT GTCACTCATG CATTGAACAA CAGCTAAGAG CAGAGCTTAG AGCTTAGAGC	540
	TGGGCCCTGT AAGGTGAGAG GAATCACATC CTGCAGAAGT CTGTCCTGAG AAGCAGGTAC	600
30	TCCTGTCACA GCAGAGACAC AGTGGATACC TGAGTAACAA TAATACAAGA CAGGACGTGG	660
	GMACAGCAAA AGATTTGGGT GTCAGAAGAR GCCGAGAACA CTTYCAGGCA GGAACATTCA	720
35	RARTTGTTCT TGGAGGAART AGGCMCSAAG GCTGGGCAGG ATTTCMCGGG GCAGAGATGG	780
	AGCAAGCAAT TGAAATGAAA GCCATGGCAT GGGAAAAGGA GCACTGGCCA CAGGGAGTGC	840
	AACGTTGTGA TGCAAGGCCA CTGTGGAGCC AT	872
40		
45	(2) INFORMATION FOR SEQ ID NO: 39:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 812 base pairs	
	(B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
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GGCAGAGGCT CACCCCAGCA GAGATTGAGG GGGAACCGTG ATGAAATTTT TAAGTATTCT 60

GCTTGATGAT AATAATTTTY CTCTTATGTT AATGITGGCT CCGTTTGGGT GTTTAGCTTT 120

TGAAAGGAGT ATGAAAATGC GGAATGGGGC TTTGGGGCTT GAGGAGGTGT GATCTCTAGT 180

60 GTTTAAAAAA TTTAATTGCA CAAATAGAAA TAATTCACCC ACATTATTGA ACCCCACTAA 240

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:

,	AGCATATCCT	TTTTGTCCAT	ATTCCTTTCC	TGCTGCCCTC	GTGTGTACCA	TTATTACTCA	300
_	GTTGTGATTT	GAGCTCGTTC	CACTTAAAGT	CATTCATAGA	TACTTTTGCG	TCGTGTTKGA	360
3	ATATTTATTG	AATTTCTATT	CTGTGTTTTA	CTTAATTACT	TTATTATGGA	ACCTTTACAC	420
	AGGTCTGGTG	TACTTGTTCT	TTGAAAAGTC	TTATGTTGAC	CACCATCACT	GAGCATATAG	480
10	CTTTTTCCTT	ATTTCCTTGG	GATAATTACC	CGAAGTGGAA	ATACCGAATC	AAACTTCTGT	540
	TTTCTTTCTT	TGGCACTATT	ATATAAATTG	TTTTCCAAAC	AAGGCATGTT	TACAATAGAC	600
15	ATTTTTCAAA	ATCTGGGTAT	TIGTCCTATT	TTGCTCTCTG	TATGCAGAAT	TCAGCGGGGT	660
15	GCCAAGTCGT	TTTCTGTGTG	GGTTGAGAGA	CAGGCTGTGC	AGCCCACTGT	TGCATAGGAC	720
	TAACTACTAC	AAATCATGCT	GAGACCGAGC	TATTTTTGCT	GCTTAGARGC	TTTGCAGCCT	780
20	TGAGTAAGTT	TCGNCATCTG	GAAACNITGN	AA			812

## 25 (2) INFORMATION FOR SEQ ID NO: 40:

30

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1515 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:

35	AATTCGGCAC (	GAGGGAAATT	CAAGCACTTT	TCCTAAAAGA	AGGGGGAATG	GATGCTGAAA	60
	CAACACGTNT (	CCACAAAGG	GAGCAGACAC	TGGGCTTGTG	AAGCTGCCCC	ATACCTTCCC	120
40	CACAGAACTG (	GCTCCGCC	TCCCTGACAT	GCAGATTTCC	ACCCAGAAGA	CAGAGAAGGA	180
40	GCCAGTGGTC 2	ATGGAATGGG	CTGGGGTCAA	AGACTGGGTG	CCTGGGAGCT	GAGGCAGCCA	240
	CCGTTTCAGC (	CTGGCCAGCC	CTCTGGACCC	CGAGGTTGGA	CCCTACTGTG	ACACACCTAC	300
45	CATGCGGACA (	CTCTTCAACC	TCCTCTGGCT	TGCCCTGGCC	TGCAGCCCTG	TTCACACTAC	360
	CCTGTCAAAG '	TCAGATGCCA	AAAAAGCCGC	CTCAAAGACG	CTGCTGGAGA	AGAGTCAGTT	420
50	TTCAGATAAG	CCGGTGCAAG	ACCGGGGTTT	CCTCCTCACG	GACCTCAAAG	CTGAGAGTGT	480
30	GGTTCTTGAG	CATCGCAGCT	ACTGCTCGGC	AAAGGCCCGG	GACAGACACT	TTGCTGGGGA	540
	TGTACTGGGC	TATGTCACTC	CATGGAACAG	CCATGGCTAC	GATGTCACCA	AGGTCTTTGG	600
55	GAGCAAGTTC	ACACAGATCT	CACCCGTCTG	GCTGCAGCTG	AAGAGACGTG	GCCGTGAGAT	660
	GTTTGAGGTC	ACGGCCTCC	ACGACGTGGA	CCAAGGGTGG	ATGCGAGCTG	TCAGGAAGCA	720
60	TGCCAAGGGC	CTGCACATAG	TGCCTCGGCT	CCTGTTTGAG	GACTGGACTT	ACGATGATTT	780
50							

	CCGGAACGTC	TTAGACAGTG	AGGATGAGAT	AGAGGAGCTG	AGCAAGACCG	TGGTCCAGGT	840
	GGCAAAGAAC	CAGCATTTCG	ATGGCTTCGT	GGTGGAGGTC	TGGAACCAGC	TGCTAAGCCA	900
5	GAAGCGCGTG	ACCGACCAGC	TGGGCATGTT	CACGCACAAG	GAGTTTGAGC	AGCTGGCCCC	960
	CGTGCTGGAT	GGTTTCAGCC	TCATGACCTA	CGACTACTCT	ACAGCGCATC	AGCCTGGCCC	1020
	TAATGCACCC	CTGTCCTGGG	TTCGAGCCTG	CGTCCAGGTC	CTGGACCCGA	AGTCCAAGTG	1080
10	GCGAAGCAAA	ATCCTCCTGG	GGCTCAACTT	CTATGGTATG	GACTACGCGA	CCTCCAAGGA	1140
	TGCCCGTGAG	CCTGTTGTCG	GGGCCAGGTA	CATCCAGACA	CTGAAGGACC	ACAGGCCCCG	1200
15	GATGGTGTGG	GACAGCCAGG	YCTCAGAGCA	CTTCTTCGAG	TACAAGAAGA	GCCGCAGTGG	1260
	GAGGCACGTC	GTCTTCTACC	CAACCCTGAA	GTCCCTGCAG	GTGCGGCTGG	AGCTGGCCCG	1320
20	GGAGCTGGGC	GTTGGGGTCT	CTATCTGGGA	GCTGGGCCAG	GGCCTGGACT	ACTTCTACGA	1380
20	CCTGCTCTAG	GTGGGCATTG	CGGCCTCCGC	GGTGGACGTG	TTCTTTTCTA	AGCCATGGAG	1440
٠	TGAGTGAGCA	GGTGTGAAAT	ACAGGCCTTC	ACTCCGTTAA	АААААААА	АААААААА	1500
25	ааааааааа	AAAAA	•			•	1515

## 30 (2) INFORMATION FOR SEQ ID NO: 41:

35

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 704 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:

40	AAGATGGTGG	CGCCCAGAGC	TTCGCTCTAT	GCTGCTCCCC	TGAGAGAGGC	GTTTCCATCA	60
	ACCAGTTTTG	CAAGGAGTTC	AATGAGAGGA	CAAAGGACAT	CAAGGAAGGC	ATTCCTCTGC	120
45	CTACCAAGAT	TTTAGTGAAG	CCTGACAGGA	CATTTGAAAT	TAAGATTGGA	CAGCCCACTG	180
70	TTTCCTACTT	CCTGAAGGCA	GCAGCTGGGA	TTGAAAAGGG	GGCCCGGCAA	ACAGGGAAAG	240
	AGGTGGCAGG	CCTGGTGACC	TTGAAGCATG	TGTATGAGAT	TGCCCGCATC	AAAGCTCAGG	300
50	ATGAGGCATT	TGCCCTGCAG	GATGTACCCC	TGTCGTCTGT	TGTCCGCTCC	ATCATCGGGT	360
	CTGCCCGTTC	TCTGGGCATT	CGCGTGGTGA	AGGACCTCAG	TTCAGAAGAG	CTTGCAGCTT	420
55	TCCAGAAGGA	ACGAGCCATC	TTCCTGGCTG	CTCAGAAGGA	GGCAGATTTG	GCTGCCCAAG	480
	AAGAAGCTGC	CAAGAAGTGA	CCCTTGCCCC	ACCAACTCCC	AGATTTCAAA	GGAGGTAGTT	540
	GCAAAAGCTG	TGCCCAAGGG	GAGGAAGGAG	GTCACACCAA	TATGATGATG	GTTTTCATGA	600
60	CTTTGAATGA	TATATTTTTG	TACATCTAGC	TGTATCGAGG	CATCAGGCCT	GAATAAACAT	660

704 

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## (2) INFORMATION FOR SEQ ID NO: 42:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1094 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

15	(xi)	SEQUENCE I	DESCRIPTION	: SEQ ID NO:	: 42:		
	GGCAGCTTTC 1	TTACAAACCC	ATCCTTCTGA	AATGTTGCTT	CAAATTCATC	CTCTGCTCCC	60
20	CAGTCCCACT	ATTCCACACA	TACTGTTACT	GTTTCTTTAT	CCTACTTTCT	CAATTTTGGA	120
20	ACATAGTTGC	AGTTACTGCA	TTGAATACCT	GTGGGTTTGC	CTGTTGTTCT	GTCTGTCTCT	180
	GTGGTTCTTG	TAATANTGGA	TCCCAGAGAT	AAAATGGACA	GTTGTNATGC	ACAGTTAATT	240
25	CAGAAACTAG	ACCTTACTTG	CTGTGTGAAA	TACCAACTAA	ATTCTCAGTG	AACTCAGCTG	300
	ANCTITATET	CCTTTTGTTT	CCCCAATTTA	TAATTTCAGT	TCAGGCCCAG	AAAGATGGAA	360
-30	TCCCAGCTAA	GAAATACAAG	TTACACCCTG	TACTAGCAGC	CCATGTGTGC	ATGTTCTTTA	420
30	AGTGCTCTTG	CAGCTATGTC	ATTTATATTG	ATTTCCCTGT	ATTATTATAA	GCAAAGCAAA	480
	TTTGAGGAAA	AAAACCCATA	ATACCACACC	TCATTTTTT	CAAGTAATAG	GGTCATAAGT	540
35	CTCATYCTYC	ATATAATATG	TTGAGTATGC	AGTATATTAT	GTGTTAGGCT	CTGGANAGGC	600
	AGAGGTTAGA	TCATGTWACA	GATÇATATCK	GATTAGGCAG	ATAAACAGTA	TTTTAACCTT	660
40	TTCCTTATTA	TATGTAACTT	GCTTTCAGGT	TTTTTAATGT	TACTATTATG	TCTTTAATAT	720
40	ATTATCTTTA	TTTGTACTTT	TGTATACAGA	GIGATTTTCC	TTTTTTAAAA	AAAATTGTGT	780
	CTTTAGGATG	GATTCCAAAG	ATGTGGAATC	AGTAGGTTTA	AGGAATATGG	ATATTTTGGC	840
45	TGGCAAGGTG	GCTCACACCT	GTAATCCCAG	CACTTTGGGA	GGCTGAGGTG	GGTGGATCAC	900
	CTGAAGTCAG	GAGTTCGAGA	CCAGCCTGAC	CAACATGGCG	AAACCCTGTT	TNTACTAAAG	960
50	ACACACWWAA	AATTRGCCAG	TGGTGGTGGC	ATGTGCTTGT	AGTCCCACTT	AGCTACTCGA	1020
JU	GAGGCTGAGG	CAGGAGAATC	GCTTGAACCC	GGGAGGCAGA	GGTTGCAGTG	AGGCAAGATG	1080
	GCACCTCTAC	ACTC					1094

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#### (2) INFORMATION FOR SEQ ID NO: 43:

60 (i) SEQUENCE CHARACTERISTICS: (A) LEXTH: 1321 base pairs (3) TYFE: nucleic acid (C) STRANDEDNESS: double

(D) TCPCLOGY: linear

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43: TOGOTTAGGO CATCACCOTT COUTTGGCTG GAACTACTGG ACAGACCCTT TTGAGATGTG 60 CCIGIGGIGC IGIGGAGATG IGIGTAGIGG TCTTAGCTCT TIGITGAGCT IGIGIGIGG 10 120 TIGISTAGIC TEASCIGIAT SCIGAAATIG GGCGTGTGTT GGAGGGCTTC TTAGCTCTTT GGTGAGATTG TATTTCTATG TGTTTGTATC ASCTGAATGT TGCTGGAAAT AAAACCTTGG 15 TTTGTMAAGG CTCYTTTTTG TGGGAAGTAA GTAGGGGAAA AGGTCTTTGA GGGTTCCTAG 300 CCTCCTTTGT ACAACAGGAA AATGCCTCAA AGCCTTGCTT CCCAGCAACC TGGGGCTGGT 360 20 TCCCAGTGCC TGGTCCTGCC CCTTCCTGGT TCTTATCTCA AGGCAGAGCT TCTGAATTTC 420 AGGCCTTCAT TCCAGAGCCC TCTTGTGGCC AGGCCTTCCT TTGCTGGAGG AAGGTACACA 480 GCGTGAAGCT GATGCTGTAC TTGGGGGATC TCCTTGGCCT GTTCCACCAA GTGAGAGAAG 540 25 GTACTTACTO TTGTACCTCC TGTTCAGCCA GGTGCATTAA CAGACCTCCC TACAGCTGTA 600 GGAACTACTG TCCCAGAGCT GAGGCAAGGG GATTTCTCAG GTCATTTGGA GAACAAGTGC 660 30 TTTAGTAGTA GTTTAAAGTA GTAACTGCTA CTGTATTTAG TGGGGTGGAA TTCAGAAGAA 720 ATTIGAAGAC CAGAICATGG GTGGTCTGCA TGTGAATGAA CAGGAATGAG CCGGACAGCC 780 TOGCTGTCAT TGCTTCTTC CTCCCCATTT GGACCCTTCT CTGCCCTTAC ATTTTTGTTT 840 35 CTCCATCTAC CACCATCCAC CAGTCTATTT ATTAACTTAG CAAGAGGACA AGTAAAGGGC 900 CCTCTTGGCT TGATTTGCT TCTTTCTTTC TGTGGAGGAT ATACTAAGTG CGACTTTGCC 960 40 CTATCCTATT TOGRAATCCC TAACAGAATT GAGTTTTCTA TTAAGGATCC AAAAAGAAAA 1020 1080 ACAAAATGCT AATGAAGCCA TCAGTCAAGG GTCACATGCC AATAAACAAT AAATTTTCCA GAAGAAATGA AATCCAACTA GACAAATAAA GTAGAGCTTA TGAAATGGTT CAGTAAGGAT 1140 45 GAGTITGITG TITTITGITT TGTTTTGTTT TGKTTTTTTA AAGACGGAGT CTCGCTCTGT 1200 CACTCAGGCT GGAGTGCAGT GGTATGATCT TGGCTCACTG TAACCTCCGC CTCCCGGGTT 1260 50 CAAGCCATTC TCCTGCCTCA GTCTCCTGAG TAGCTGGGAT TACAGGTGCG TGCCACCATG 1320 CCTGGCTAAT TITTGTGTTT TTAGTAGAGA CAGGGTTTCA CCATGTTGGT CGGGCTGGTC 1380 TCAAACTCCT GACCTCTTGA TCCGCCTGCC TTGGCCTCCC AAAGTGATGG GATTACAGAT 1440 55 GTGAGCCACC CGTSCCCTAG CCAAGGATGA GATTTTTAAA GTATGTTTCA GTTCTGTGTC 1500 ATGGTTGGAA GACAGAGTAG GAAGGATATG GAAAAGGTCA TGGGGAAGCA GAGGTGATTC 1560 60 ATGGCTCTGT GAATTTGAGG TGAATGGTTC CTTATTGTCT AGGCCACTTG TGAAGAATAT 1620 WO 98/54963 PCT/US98/11422

	GAGTCAGTTA TTGCCAGCCT TGGAATTTAC TTCTCTAGCT TACAATGGAC CTTTTGAACT	1680
5	GGAAAACACC TTGTCTGCAT TCACTTTAAA ATGTCAAAAC TAATTTTTAT AATAAATGTT	1740
3	TATTTTCACA TTGAAAAAAA AAAAAAATTT AAAAACYCGG GGGGGCCCCS GHACCCCATT	1800
	NGCCCCTAAG GGGGGGGTT T	1821
10		
	(2) INFORMATION FOR SEQ ID NO: 44:	
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1024 base pairs	
•	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:	
	GGGGCACAGT TGAAGAAGCG ACCGAGGGAC TGGGAGTCGT TAGTGAGGAT GACGCGGCAT	60
25	GGCAAGAACT GCACCGCAGG GCCGTCTACA CCTACCACGA GAAGAAGAAG GACACAGCGG	120
	CCTCGGGCTA TGGGACCCAG AACATTCGAC TGAGCCGGGA TGCCGTGAAG GACTTCGACT	180
20	GCTGTTGTCT CTCCCTGCAG CCTTGCCACG ATCCTGTTGT CACCCCAGAT GGCTACCTGT	240
30	ATGAGCGTGA GGCCATCCTG GAGTACATTC TGCACCAGAA GAAGGAGATT GCCCGGCAGA	300
	TGAAGGCCTA CGAGAAGCAG CGGGGCACCC GGCGCGAGGA GCAGAAGGAG CTTCAGCGGG	360
35	CGGCCTCGCA GGACCATGTG CGGGGCTTCC TGGAGAAGGA GTCGGCTATC GTGAGCCGGC	420
	CCCTCAACCC TTTCACAGCC AAGGCCCTCT CGGGCACCAG CCCAGATGAT GTCCAACCTG	480
40	GGCCCAGTGT GGGTCCTCCA AGTAAGGACA AGGACAAAGT GCTGCCCAGC TTCTGGATCC	540
40	CGTCGCTGAC GCCCGAAGCC AAGGCCACCA AGCTGGAGAA GCCGTCCCGC ACGGTGACCT	600
	GCCCCATGTC AGGGAAGCCC CTGCGCATGT CGGACCTGAC GCCCGTGCAC TTCACACCGC	660
45	TAGACAGCTC CGTGGACCGC GTGGGGCTCA TCACCCGCAG CGAGCGCTAC GTGTGTGCCG	720
	TGACCCGCGA CAGCCTGAGC AACGCCACCC CCTGCGCTGT GCTGCGGCCC TCTGGGGCTG	780
50	TGGTCACCCT CGAATGCGTG GAGAAGCTGA TTCGGAAGGA CATGGTGGAC CCTGTGACTG	840
50	GAGACAAACT CACAGACCGC GACATCATCG TGCTGCAGCG GGGCGGTACC GSTTCGCGGG	900
	CTCCGGAGTG AAGCTGCAAG CGGAGAAATC ACGGCCGGTG ATGCAGGCCT GAGTGTGTGC	960

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(2) INFORMATION	FOR	SEQ	ID	NO:	45:
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1:1	CENTENICE	CHARACTERISTICS:
(1)	SECUENCE	CHARACTERISTICS:

(A) LENGTH: 983 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10	(xi)	SEQUENCE D	ESCRIPTION:	SEQ ID NO:	45:		
	CGACACGGCT G	CGAGAAGAC (	GACAGAAGGG	CCCGACCGCG	AGCCGTCCAG	GTCTCAGTGC	60
15	TGTGCCCCCC C	CAGAGCCTA	GAGGATGTTT	CATGGGATCC	CAGCCACGCC	GGGCATAGGA	120
IJ	GCCCTGGGA A	CAAGCCGGA	GCTGTATGAG	GAAGTGAAGT	TGTACAAGAA	CGCCCGGGAG	180
	AGGGAGAAGT A	CGACAACAT	GGCAGAGCTG	TTTGCGGTGG	TGAAGACAAT	GCAAGCCCTG	240
20	GAGAAGGCCT A	CATCAAGGA	CTGTGTCTCC	CCCAGCGAGT	ACACTGCAGC	CTGCTCCCGG	300
	CTCCTGGTCC A	ATACAAAGC	TGCCTTCAGG	CAGGTCCAGG	GCTCAGAAAT	CAGCTCTATT	360
25	GACGAATTCT C	SCCGCAAGTT	CCGCCTGGAC	TGCCCGCTGG	CCATGGAGCG	GATCAAGGAG	420
23	GACCGGCCCA T	CACCATCAA	GGACGACAAG	GGCAACCTCA	ACCGCTGCAT	CGCAGACGTG	480
	GTCTCGCTCT 1	CATCACGGT	CATGGACAAG	CTGCGCCTGG	AGATCCGCGC	CATGGATGAG	540
30	ATCCAGCCCG A	ACCTGCGAGA	GCTGATGGAG	ACCATGCACC	GCATGAGCCA	CCTCCCACCC	606
	GACTTTGAGG (	GCCGCCAGAC	GGTCAGCCAG	TGGCTGCAGA	CCCTGAGCGG	CATGTCGGCG	660
35	TCAGATGAGC T	rggacgactc	ACAGGTGCGT	CAGATGCTGT	TCGACCTGGA	GTCAGCCTAC	72
	AACGCCTTCA A	ACCGCTTCCT	GCATGCCTGA	GCCCGGGGCA	CTAGCCCTTG	CACAGAAGGG	. 78
	CAGAGTCTGA (	GCGATGGCT	CCTGGTCCCC	TGTCCGCCAC	ACAGGCCGTG	GTCATCCACA	84
40	CAACTCACTG	CTGCAGCTG	CCTGTCTGGT	GTCTGTCTTT	GGTGTCAGAA	CTTTTGGGCC	90
	GGGCCCCTCC (	CCACAATAAA	GATGCTCTCC	GACCTTCAAA	АААААААА	AAAAAAAAGR	96
45	KGSGGCCGGT	CCCCANTCCC	ccc				98

#### (2) INFORMATION FOR SEQ ID NO: 46:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2421 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:

CCGGCTGATC GCTGCCGCTC CGCCAATACA ATAGAGCCAK CCACTACCAG CAGCCTGGCC

60

60

50

	CTCTTCCTCC	TTCTCCAGAG	AGACCAATCC	AGCCGAACTC	GGGGTTTGCC	TGAGGAGAAG	120
	GAGGAAGTGA	CCATGGACAC	AAGTGAAAAC	AGACCTGAAA	ATGATGTTCC	AGAACCTCCC	180
5	ATGCCTATTG	CAGACCAAGT	CAGCAATGAT	GACCGCCCGG	AGGGCAGTGT	TGAAGATGAG	240
	GAGAAGAAAG	AGAGCTCGCT	GCCCAAATCA	TTCAAGAGGA	AGATCTCCGT	TGTCTCAGCT	300
10	ACCAAGGGGG	TGCCAGCTGG	AAACAGTGAC	ACAGAGGGGG	GCCAÇCCTGG	TCGGAAACGA	. 360
10	CGCTGGGGAG	CCAGCACAGC	CACCACACAG	AAGAAACCTT	CCATCAGTAT	CACCACTGAA	420
	TCACTAAAGA	GCCTCATCCC	CGACATCAAA	CCCCTGGCGG	GGCAGGAGGC	TGTTGTGGAT	480
15	CTTCATGCTG	ATGACTCTCG	CATCTCTGAG	GATGAGACAG	AGCGTAATGG	CGATGATGGG	540
	ACCCATGACA	AGGGGCTGAA	AATATGCCGG	ACAGTCACTC	AGGTAGTACC	TGCAGAGGGC	600
20	CAGGAGAATG	GGCAGAGGGA	AGAAGAGGAA	GAAGAGAAGG	AACCTGAAGC	AGAACCTCCT	660
20	GTACCTCCCC	AGGTGTCAGT	AGAGGTGGCC	TTGCCCCCAC	CTGCAGAGCA	TGAAGTAAAG	720
	AAAGTGACTT	TAGGAGATAC	CTTAACTCGA	CGTTCCATTA	GCCAGCAGAA	GTCCGGAGTT	780
25	TCCATTACCA	TTGATGACCC	AGTCCGAACT	GCCCAGGTGC	CCTCCCCACC	CCGGGGCAAG	840
	ATTAGCAACA	TTGTCCATAT	CTCCAATTTG	GTCCGTCCTT	TCACTTTAGG	CCAGCTAAAG	900
30	GAGTTGTTGG	GGCGCACAGG	AACCTTGGTG	GAAGAGGCCT	TCTGGATTGA	CAAGATCAAA	960
50	TCTCATTGCT	TTGTAACGTA	CTCAACAGTA	GAGGAAGCTG	TTGCCACCCG	CACAGCTCTG	1020
	CACGGGGTCA	AATGGCCCCA	GTCCAATCCC	AAATTCCTTT	GTGCTGACTA	TGCCGAGCAA	1086
35	GATGAGCTGG	ATTATCACCG	AGGCCTCTTG	GTGGACCGTC	CCTCTGAAAC	TAAGACAGAG	1140
	GAGCAGGGAA	TACCACGGCC	CCTGCACCCC	CCACCCCCAC	CCCCGGTCCA	GCCACCACAG	120
40	CACCCCCGG	CAGAGCAGCG	GGAGCAGGAA	CGGGCAGTGC	GGGAACAGTG	GGCAGAACGG	126
. •	GAACGGGAAA	TGGAGCGGCG	GGAGCGGACT	CGATCAGAGC	GTGAATGGGA	TCGGGACAAA	132
	GTTCGAGAAG	GCCCCCTTC	CCGATCAAGG	TCCCGTRACC	GCCGCCGCAA	GGAACGTGCG	138
45	AAGTCTAAAG	AAAAGAAGAG	TGAGAAGAAA	GAGAAAGCCC	AGGAGGAACC	ACCTGCCAAG	144
	CTGCTGGATC	ACCITITCCC	AAAGACCAAG	GCAGCTCCCI	GCATCTATTG	GCTCCCACTG	150
50	ACTGACAGCO	AGATCGTTCA	A GAAAGAGGCA	A GAGCGGGCCG	AACGGGCCAA	GGAGCGGGAG	156
	AAGCGGCGAA	AGGAGCAAGA	A AGAAGAAGAC	CAAAAGGAGG	GGGAGAAGGA	AGCCGAGCGG	162
	GAACGGAAC	C GACAGCTGG	GCGAGAGAAA	A CGTCGGGAGG	: ACAGTCGGGA	GAGGGACAGG	168
55	GAGAGAGAG	A GAGAAAGGG/	A GCGGGACAGO	GGGGACCGAC	ATCGGGATAC	GGAAAGGGAC	174
	CGAGAACGAC	GCAGGGAAAG	GGATCGCAG	GACACCAAGO	GCCACAGCAC	AAGCCGGAGT	180
60	CGGAGCACAG	CTGTGCGGG	A CCGGGGTGGG	GCCGCTAGG	TGGGAAAACA	CTAGAGCTGC	186

WO 98/54963

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•	AGGTACCAGC CACTCGGCCC CAGGGGGTTA TGGCCACAGA GGGATAGGCA CAGTCTCCAC	1920
÷	CACCCTGGAG CCAAGGGTCT TTCACATCAC CTATCCCTAC ATACATACCA AATGGAAAAG	1930
5	TGGCCATCCT TTTCCCCCCA AACACACCCC CTTAACCTAT CTCTTGGGAC TTAGCCCGAC	2340
	CCTCCCTCTC ATTTCCCATT AAGTCTGAGA GGCAAGAGCT AGGTTAGGCA AGGAGGTGGT	2100
	TGGCCAGAGA TGGGGAACAG CCAGGTGCCC CAGTCCTCTG ATTITTCCTC CATCCTGCTT	2160
10	ACCACCTCCC TGGGTACTTA CAGCCTTCTC TTGGGAACAG CCGGGGCCAG GACTGGGTCA	2220
	CCTATGAGCT GAATCAGCAT CTCCTCCTGA GTCCCAGGGC CCCTGCAGTT CCCAGTCTCT	2280
15	TCTGTCCTGC AGCCCTTGCC TCTTTCCCAC AGGTTCCACT TTATATCCAC CTTTTCCTTT	2340
	TGTTCAATTT TTATTTTAT TTTTTTTATT ATTAAATGAT GTGGTCTATG GAAAAAAAA	2400
20	TAAAAATCTG ACTTAGTFFF A	2421
20		
	(2) THEODAY TOD GEO TO NO. 47.	
25	(2) INFORMATION FOR SEQ ID NO: 47:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 840 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
30	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:	
35	CTCAAACTCC TGAGCTGAAG CGATCTACCT GCCTCAGCTA GGATTACAGG TGTGAGCCAC	60
	CGCACCCAAC CTCAATAAGC KTATTTGATA AAAKATATGC AAGCTCCCTT TATKCACTTT	120
	TCATTCAGAA TGTTTAGTAA TTTGTATTGT TTTTCAGATT TTCAGCCCAA TATATCTCCY	180
40	TGCCCACTGT GTCACTGTAT TCTACCTAWA CATCATCACG TGTTTCTGCT ATTGGCTGTA	240
	TGATGGAACA CTGCGGCTCA TTTTCCTGAA AACTGCCGAT AGTGCATAGA RTGCTGGGAT	300
45	GGAAACCAGA ARCTTTGAAT TCAAGCCTTG GTTCTGCCTT GTTTTTGCTT GGGTGGCCTT	360
43	GAGTCAGCCA CATACCTTTT AAAATCTCAA TTTATTAGAA ATTATTCCAA ATCAAAATCA	420
	AATGAGAAGG TATATACAAA AGTGCTTTAT CCCACAATAA ACTATTCAAG AGAGAGCAAA	180
50	GGAGAGGACA TTTACTCAAC ACCTCCTAAA AGGCAGCCAG TGAAATTAGG CATTTTATTT	540
	AATCCTCCTG GCAACTCTGA GAGTAAAGCA TTATTAATCC CATTTTGGCT GTTTAAAGAA	500
<b>5</b>	ATTATTTGCA CTAGATTCCA GCTGTAGTTT AGYTTCAGAA AAAAAAATCC TGAGATGTGA	56
55	ATTCACAGCT TTCTGGGTTT AAAGCCCAAG CTCTATCACA TCATGCTATT ATTGTTACAT	72
	TACTGCTAGT TCTATGAAAA GAAATACTAA TTTATGAAAT ACATCTTATC CAAAAAAAAA	78

AAAAAAAAC TGGGAGGGG GGCCCGTACC CAAATCGCCG GATAGTGATC GTAAACAATC

5 (2)	INFORMATION	FOR	SEQ	ID	NO:	48:
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2432 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:

		_					
15	GGCACGAGGC	CCGCAACGCT	GAGGAAGGGC	CCGTCCCGCC	TTCCCCGGCG	CGCCATGGAG	60
	ccccccccc	TTGCAGAAGC	CGTGGAGACG	GGTGAGGAGG	ATGTGATTAT	GGAAGCTCTG	120
20	CGGTCATACA	ACCAGGAGCA	CTCCCAGAGC	TTCACGTTTG	ATGATGCCCA	ACAGGAGGAC	180
20	CGGAAGAGAC	TGGCGGASTG	CTGGTCTCCG	TCCTGGAACA	GGGCTTGCCA	CCCTCCCACC	240
	GTGTCATCTG	GCTGCAGAGT	GTCCGAATCC	TGTCCCGGGA	CCGCAACTGC	CTGGACCCGT	300
25	TCACCAGCCG	CCAGAGCCTG	CAGGCAYTAG	CCTGYTATGY	TGACATCTCT	GTCTCTGAGG	360
	GGTCCGTCCC	AGAGTCCGCA	GACATGGATG	TTGTACTGGA	GTCCCTCAAG	TGCCTGTGCA	420
30	ACCTCGTGCT	CAGCAGCCCT	GTGGCACAGA	TGCTGGCAGC	AGAGGCCCGC	CTAGTGGTGA	480
30	AGCTCACAGA	CCTCTCCGG	CTGTACCGTG	AGAGGAGCTT	CCCCACGAT	GTCCAGTTCT	540
	TTGACTTGCG	GCTCCTCTTC	CTGCTAACGG	CACTCCGCAC	CGATGTGCGC	CANAGCTGTT	600
35	TCAGGAGCTG	AAAGGAGTGC	GCCTGCTAAC	TGACACACTG	GAGCTGACGC	TGGGGGTGAC	660
	TCCTGAAGGG	AACCCCCAC	CCACGCTCCT	TCCTTCCCAA	GAGACTGAGC	GGGCCATGGA	720
40	GATCCTCAAA	GIGCTCTTCA	ACATCACCCT	GGACTCCATC	AAGGGGGAGG	TGGACGAGGA	780
40	AGACGCTGCC	: CTTTACCGAC	: ACCTGGGGAC	CCTTCTCCGG	CACTGTGTGA	TGATCGCTAC	840
	TGCTGGAGAC	: CGCACAGAGG	AGTTCCACGG	CCACGCAGTA	ASCCTCCTGG	GGAACTTGCC	900
45	CCTCAAGTGT	CTGGATGTTC	TCCTCACCCT	GGAGCCACAT	GGAGACTCCA	CGGAGTTCAT	960
	GGGAGTGAAT	· ATGGATGTGA	TTCGTGCCCT	CCTCATCTTC	CTAGAGAAGC	GTTTGCACAA	1020
50	GACACACAGO	G CTGAAGGAGA	GTGTAGCTCC	CGTGCTGAGG	GTGCTGACTG	AATGTGCCCG	1080
30	GATGCACCGC	CCAGCCAGGA	AGTTCCTGA	GCCCAGGTC	CTGCCCCCTC	TGCGGGATGT	1140
	GAGGACACG	CCTGAGGTTC	GGGAGATGC	C GCGGAACAAC	CTTGTCCGCC	TCATGACACA	1200
55	CCTGGACAC	A GATGTGAAG	A GGGTGGCTG	CGAGTTCTTC	TTTGTCCTG	GCTCTGAGAG	1260
	TGTGCCCCG	A TTCATCAAGT	r ACACAGGCT	A TGGGAATGC	CTGCCTTC	TGGCTGCCAG	1320
60	GGGCCTCATY	G GCAGGAGGC	G GCCCGAGGG	C AGTACTCAGA	A GGATGAGGAC	: ACAGACACAG	1380
υU							

•	ATGAGTACAA	GGAAGCCAAA	GCCAGCATAA	ACCCTGTGAC	CGGGAGGGTG	GAGGAGAAGC	1440
	CGCCTAACCC	TATGGAGGGC	ATGACAGAGG	AGCAGAAGGA	GCACGAGGCC	ATGAAGCTGG	1500
5	TGACCATGTT	TGACAAGCTC	TCCAGGAACA	GAGTCATCCA	GCCAATGGGG	ATGAGTCCCC	1560
	GGGTCATCT	TACGTCCCTG	.CAGGATGCCA	TGTGCGAGAC	TATGGAGCAG	CAGCTCTCCT	1620
10	CGGACCCTGA	CTCGGACCCT	GACTGAGGAT	GGCAGCTCTT	CTGCTCCCCC	ATCAGGACTG	1680
LO.	GTGCTGCTTC	CAGAGACTTC	CTTGGGGTTG	CAACCTGGGG	AAGCCACATC	CCACTGGATC	1740
	CACACCCGCC	ÇCCACTTCTC	CATCTTAGAA	ACCCCTTCTC	TTGACTCCCG	TTCTGTTCAT	1800
15	GATTTGCCTC	TGGTCCAGTT	TCTCATCTCT	GGACTGCAAC	GGTCTTCTTG	TGCTAGAACT	1860
	CAGGCTCAGC	CTCGAATTCC	ACAGACGAAG	TACTTTCTTT	TGTCTGCGCC	AAGAGGAATG	1920
20	TGTTCAGAAG	CTGCTGCCTG	AGGGCAGGGC	CTACCTGGGC	ACACAGAAGA	GCATATGGGA	1980
20	GGGCAGGGGT	TTGGGTGTGG	GTGCACACAA	AGCAAGCACC	ATCTGGGATT	GGCACACTGG	2040
	CAGAGCMANT	GTKTTGGGGT	ATGTGCTGCA	CTTCCCAGGG	AGAAAACCTG	TCAGAACTTT	2100
25	CCATACGAGT	ATATCAGAAC	ACACCCTTCC	AAGGTATGTA	TGCTCTGTTG	TTCCTGTCCT	2160
	GTCTTCACTG	AGCGCAGGGC	TGGAGGCCTC	TTAGACATTC	TCCTTGGTCC	TCGTTCAGCT	2220
30	GCCCACTGTA	GTATCCACAG	TGCCCGAGTT	CTCGCTGGTT	TTGGCAATTA	AACCTCCTTC	2280
50	CTACTGGTTT	AGÁCTACACT	TACAACAAGG	AAAATGCCCC	TCGTGTGACC	ATAGATTGAG	2340
	ATTTATACCA	CATACCACAC	ATAGCCACAG	AAACATCATC	TTGAAATAAA	GAAGAGTTTT	2400
35	GGACAAAAA	АААААААА	ааааааааа	AA	•		2432

## 40 (2) INFORMATION FOR SEQ ID NO: 49:

45

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1742 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:

50	GTCCTGCAGG	AGCTGCACGC	GCCCGAGGTG	CGCANGAACA	AGGAGCAGCG	AGAAGAGATG	60
	TCGGGCTAAG	GGCCCGGSAC	GRGSGGCGCC	CATCCTGCGA	CGGAACACGT	TCGGGTTTTG	120
55	GTTTTGTTTC	GTTCACCTCT	GTCTAGATGC	AACTTTTGTT	CCTCCTCCCC	CACCCCAGCC	180
33	CCCAGCTTCA	TGCTTCTCTT	CCGCACTCAG	CCGCCCTGCC	CTGTCCTCGT	GGTGAGTCGC	240
	TGACCACGGC	TTCCCCTGCA	GGAGCCGCCG	GGCGTGRAGA	CGCGGTCCCT	CGGTGCAGAC	300
60	ACCAGGCCGG	GCGCGGCTCG	GTCCCCCGGG	GGCCCTGTGA	GAGAGGTGGY	GGTGACCGTG	360

	GTAAACCCAG	GGCGGTGGCG	TGGGATCRCG	GGTCCTTACG	CTGGGCTGTC	TGGTCAGCAC	· <b>4</b> 20
5	GTGCAGGTCA	GGGCAGGTCC	TCTGAGCCGG	CGCCCCTGGC	CAGCAGGCGA	GGCTACAGTA	480
3	CCTGCTGTCT	TTCCAGGGGG	AAGGGGCTCC	CCATGAGGRA	GGGGCGACGG	GCGAGGGGGG	540
	TGATGGTGCC	TGGGAAGCCT	GCKTGTGCAN	CCGGTGCTTG	TTGAACTGGC	AGGCGGGTGG	600
0	GTGGGGGCTG	CAGCTTTCCT	TAATGTGGTT	GCACAGGGGT	CCTCTRAGAC	CACCTGGCGT	660
	GAGGTGGACA	CCCTGGGCCT	TCCTGGAAGC	CTGCAGTTGG	GGGCCTGCCC	TGAGTCTGCT	720
.5	GGGGAGTGGG	CATTCTCTGC	CAGGGACCCA	TGAGCAGGCT	GCATGGTCTA	GAGGTTGTGG	780
	GCAGCATGGA	CAGTCCCCCA	CTCAGAAGTG	CAAGAGTTCC	AAAGAGCCTC	TGGCCCAGGC	840
	CCCTCCCTGG	GACAGCCCCG	CCGCCCTCC	CCACCAGGGC	TTTGCAGATG	TCCTTGAAAG	900
20	ACCCACCCTA	GAGCCCTTTG	GAGTGCTGGC	CCCTCCTGTG	CCCTCTGCCC	TGGTGGAAGC	960
	GGCASCACAA	GTCCTCCTCA	GGGAGCCCCA	AGGGGGATTT	TKTGGGACCG	CTGCCCACAG	1020
25	ATCCAGGTGT	TGGAAGGGCA	GCGGGTAAGG	TTCCCAAGCC	AGCCCCAACA	CCCTTCCCAC	1080
	TTGGCACCCA	GAGGGGGCTG	TGGGTGGAGG	CCTGACTCCA	GCCTCTCCT	GCCCACACCC	1140
	TCTGGGCTGA	GTTCCTTCTT	TCCCTTGGAC	GCCCAGTGCT	GCCTTGGAG	GACGGTCAGC	1200
30	TGGAGGATGG	CGGTGGGGGA	GGCTGTCTTT	GTACCACTGC	AGCATCCCCC	ACTTCTCCAC	1260
	GGAAGCCCCA	TCCCAAAGCT	GCTCCCTCCC	CCCTTGCTGI	' AAAGTGTGAA	GGGGGGGCT	1320
35	GAGTTCTCTT	AGGACCCAGA	GCCAGGGCCC	TCAACTTCCA	TCCTGCGGGA	GGCCTTGGCC	1380
55	GGGCACTGCC	AGTGTCTTCC	AGAGCCACAC	CCAGGGACCA	CGGGAGGATC	CTGACCCCTG	1440
	CAGGGCTCAG	GGGTCAGCAG	GGACCCACTG	CCCCATCTCC	CTCTCCCCAC	CAAGACAGCC	1500
40	CCAGAAGGAG	CAGCCAGCTG	GGATGGGAAC	: CCAAGGCTGT	CCACATCIGG	CTTTTGTGGG	1560
	ACTCAGAAAG	GGAAGCAGAA	CTGAGGGCTC	GGATATICCI	CATGGTGGCA	GCGCTCATAG	1620
45	CGAAAGCCTA	CTGTAATATG	CACCCATCTC	ATCCACGTAC	TAAAGTGAAC	TTAAAAATTC	1680
1.5	AATCAAATGA	ACAATTAAAT	AAACACCTGT	GTGTTTAAG	AAAAAAAA A	AAAAAAACTG	1740
•	CG		•				1742

#### (2) INFORMATION FOR SEQ ID NO: 50:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1487 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(XI)	SEQUENCE	DESCRIPTION:	SEQ	TD	NO:	50

	GGCACGAGCC	TCCGCGAACT	GTGGAGTCGG	CGGAGGGCTG	GAATCAGCGT	GGGCTCCAGG	60
5	TCGCTGGCAG	CCGGGTGGCA	GAACTCTTCC	GAGGCTCCTT	GGGAAGAAGC	TACACCCGAG	120
	GGAGCCGGAT	GGGCCTCGAA	AACCTGGCCC	GCTCTGGTTC	TGTACCATTG	CAAGGGGAAC	180
	CGTAAACTGA	GCTTTTCTAA	CGTGGGTTTC	TGCCAAGTAC	TTTTCCAGCT	GCCCCCTTCC	240
10	CCCCAGCACA	CAGGAGAGCC	TCTGTGTAGC	CAGCGCTTGA	CAGTCGTTAG	GTAGGTTGTA	300
	CTGTGTAGGG	AGGAGCTCAA	GATCATGAAT	GGTTGTCACA	GGAGAAAGCG	GTTGCATCTT	360
15 .	TGCAAAACTA	TATACCTGCT	CTCCTTTCTC	TTTTCTTTTC	TGCTGAGTAA	TGAAGTTGTA	420
	AGTTCACACT	GGCACATTCT	CAGGGCTGTG	CAGATTATTT	GCACTTTATT	TCATAGGTGR	480
20	ATAAGTGCTT	TTTAGCTTTC	TTTGTATATT	GAGTTGCTTT	TGAATTGCTT	CCCATATTTT	540
20	TATTTCATAC	AAACTGAACA	ATTGTGGCCC	CTCTATTTTA	TTTATAAAGG	TTCAGTGTAT	600
	CTTTGCCTGC	CTACATCAAT	CTGCAAGGGA	GTTGCAGAAA	GCCTCATGTT	CATCGAGCCG	660
25	TGAGTCACAA	CCAATTTCTA	AGCTGTTATA	ACAAAAAAGT	GTTTGCTTTT	TTTCACAAGT	720
	AACTTTAAAA	GTGTAGTTTA	GAAAGAAAAC	ATTTTCAATA	AAAAGACACT	ACATTAATCC	780
30	TGGATGCTTG	CAAATCCTAA	AATMTATTCC	TCCTCTAGCG	TTGCACAGCT	CTGTGTTGTA	840
50	TACACAGACT	AGCTTTAAAA	TTTGTCACAT	ACCACTTTAC	CTTTACTTTT	ATGTATCATT	900
	CCCCCGACTT	CCTTACTGCA	GGTGTGGGCA	AGAAAACTTT	TCCTTTAACA	CTTTTCAACA	960
35	GCGGGCATAA	AATTCTGCAG	CTGAGGTCTT	GAAGAATGCA	GATGGGTACA	GTATGTGTTG	1020
	GAGCTCACAG	TGTGTATTGA	CTAACCTAGT	TCCTTTTTTG	CTTTTTTTGG	TATTGTCTTG	1080
40	TTAAAAGTGA	CTCCCAGGTA	GCAACTCTCT	TTTTTAAGGG	TGGGAACGAA	AGGGACGTAG	1140
10	GAAGAATAGA	TCTAGATTAT	TTAACAGTCT	TCGATAGAGT	TTGAAAGCTT	TCTTCTTCAT	1200
	TCAATTTTGG	GCAAAATACT	GCCTCTGCAT	TTGTTCATAA	CAAAAAGATT	AGATTAATAA	· 1260
45	GTAGCTTTTG	TTGGTGGAAA	TTACCAGCTC	TATAAGTCAC	CCTTGGTGGT	TCATGGACCT	1320
	CTGATTAGCT	TGGGTTTTGC	AGTCTCATTG	CCACATGTAT	ATGTGGAGCC	AATGGCCTTT	1380
50	TGGTGCTCAG	CTGTTTACGT	CTGACTCCTT	GACTTCTTTG	GTACAGTGAT	GGAGTCAGAT	1440
<i>-</i>	CTCATTAAGT	GTGATTCTCC	ATGGATATAA	CCAGCCCCAA	AAAAANG		148

- (2) INFORMATION FOR SEQ ID NO: 51:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 1328 base pairs

60 (B) TYPE: nucleic acid

(C) STRANDEDNESS: double(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 51:

5			
J	GGCACGAGCT CGTGCCGAAT TCGGCACGAG AGAAGAT	TG AAGAAGCCAG ATCCAC	GCTTC 60
	CCTGCGGGCT GCTTCTTGTG GGGAAGGGAA AAAGAGG	AG GCCTGTAAGA ACTGC	ACCTG 120
10	TGGCCTTGCC GAAGAACTGG AAAAAGAGAA GTCAAGGC	BAA CAGATGAGCT CCCAAG	CCCAA 180
	GTCAGCTTGT GGAAACTGCT ACCTGGGCGA TGCCTTCC	CGC TGTGCCAGCT GCCCC	TACCT 240
15	TGGGATGCCA GCCTTCAAAC CTGGGGAAAA GGTGCTTG	TG AGTGATAGCA ATCTT	CATGA 300
13	TGCCTAGGAG GTTCCTGACA TGGGACCCAT CTGCTCC	PCC AGCCAACTCC TGTCC	CTCAC 360
٠	ATCCCACCAT GGTGGCTCCT CCCACCTCCT CTGGATT	IGT TCACTCTGAG ATCTG	rttgc 420
20	AGAGTGGGTG CTTAGCAGAC AGAGTGAAGC TGGCTGG	egg gcacagtegt eteta	GTGCT 480
	GCTGTGTATC AAAAGACCAA GGTATTATGG GACCTGG	ITT CAGAATGGGA TGGGT	TTCTT 540
25	CACCTCATGT TAAGAGAAGG GAGTGTGTCC TGAAGAA	GCC CTTCTTCTGA TGTTA	OOO STAAA
25	CTGACCAGAA CGCTCTTGAG CCCAGGCATC GTTGAGC	ATT AACACTCTGT GACAG	AGCTG 660
	CAGACCCCTG CCTTGAGTCT CATCTCAGCA ATGCTGC	CAC CCTCTTGTCT TICAG	AGTTG 720
30	TTAGTTTACT CCATTCTTTG TGACACGAGT CAAGTGG	CTC ACAACCTCCT CAGGG	CACCA 780
	GAGGACTCAC TCACTGGTTG CTGTGATGAT ATCCAGT	GTC CCTCTGCCCC CTTCC	ATCCC 840
35	CAACCACATT TGACTGTAGC ATTGCATCTG TGTCCTG	TTG TCATTTATGT TAACC	TTCAG 900
دد	GTATTAAACT TGCTGCATAT CTTGACATAT CTTGAGA	TTC TGCATGTCTT GTAAA	GAGAG 960
	GGGATGTGCA TTTGTGTGTG ATGTTGGATA GTCATCO	ACG CTCAGTTTGG ACCAT	TGGAG 1020
40	GAACTTAGTG TCACGCACAA ATGGGGCTAT TCCTACG	CTT AGAATAGGGC TTGTC	TGCCC 1080
٠	ACTITAGAAG AGTCCCAGGT TGGTGAGCAT TTAGAGG	GAA GCAGGGCAGA ACTCI	GAACG 1140
45	ACAATACGTC TCTCTGAGCA GAGACCCCTT TGTTCTT	GTT ATCCACCCAT ATGG	CTTGG 1200
40	AATCAATCTT GCCAAATATT TGGAGAGATT GTGTGGA	TTT AAGAGACCTG GATTI	<b>TTATA</b> 1260
	TTTTACCAGT AAATAAAAGT TTTCATTGAT ATCTGTC	CTT GAAAAAAAA AAAA	1320 AAAAA
50	AAACTCGA		1328

55 (2) INFORMATION FOR SEQ ID NO: 52:

60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1856 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

## (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ □ NO: 52:

	(XI) SEQUENCE LESCRIFICATION, SEQ LO NO. 34.	
5	GAATTCGGCA CGAGCTCTGC AACATTGCAA ATGAACTTGT AGGCGAGGGT TCCGCTGCCC	60
	CCTAGATTAA ATTCCCCGGG CTGAAACTGA GTTGCAGATT TACAATATCA TATTTTAAAT	120
10	TGCTGTCTTC AATTAAACCA TTTATGACCA TAACTAATTT TCAGGATGTC GATGCATGCT	180
10	TITTCCAGGCC TITCCTTCTTT GTACAAAAGT AAATGTCCAT AAAGCGTTTC ACTTATATTC	240
	TTCAAACATG ATGCTAATTT AAATTAATTA CTTCCTATGA TATGTTATTA TTCCTATGAT	300
15	TTTGCCACTG TTATTAGTTC TCTCAAAAAT ACATCTAGGG AAGAGGATTA TTTTAAGTRA	360
	TTTGATTATC TTTCTATCTC TTTTATTTAT TTCTCATTTA CITAAGAAAT TCGTTCCATT	420
20	GGTTGGCATT GATACAGTAA ATTTGTAAAT GAGGAGACAA TATAAAAAAT CTAAATTACT	480
	TGTGCTTAAT GACIGTAGCA GAATSCCTTT TCTCTAAATC AGAITGTCTT TCTTGCAGTT	540
	TAGTITGATA GATITGCAAG CTATGCTSCT TCCATGAAGT TAGCTSCGCT GGTAGGAACG	600
25	CAGGCTTCTT TGTCTCTGGT TGTAGCTTGC ATGATCGCCC CATTAGGCAG ACAACGTAGC	660
	CGGAGATCAC AAATCAGGCC CTTGGTGTAG TTGCTAGTGT GTGGAGGTGC AGAGAGGTTG	720
30	GCAGAAACTG ACCTCACTGG GCAAGGGTGG CCATGGACCT GAITCTTTAA TGCACTCTAT	780
50,	GTGTTCAGGA AGCCACAGGC CATATTTEAT TOTGAGAAAA AAAALAAGAG GAAAAACCCC	840
	ACAAAGTATA ACAACCOCTT AAGATACATC TATYYTAAAS TGAAATTAAT TTTTCAGTTT	900
35	ATACCATTGG CCAATTACAA GATAAAAATG TTCAATTTCT TIAAGAATCC TTTGTTGACT	960
	TGTCTTTTCA TCTCTTGCTA TTTATATUTG TCACTGTTAG TCAACAAAGT CTTATTTGCT	1020
40	GAGGAAGGAC TITGCTGCAC TTACTGTACT ACATCAAACA CTGGGGAGGG TGGTGTTTAA	1080
	CTTTTTAAAA AATGTTATTC TGATTATAAD AATAATATTG GCTTTTTTCA TGAAAAGAGC	1140
	GCCACCTTGC AAGGTTTAGT GAGATTTATG GAAGTTGAAT ACCTAAGCAG GAATTGCTGC	1200
45	TAGCTCCAAA AATTTGCGAA GCAAAASCTA GCCCCAATTS STFTGGAAGT TTGAAACTGA	1260
	TTAACAGATT TGCATTTGAA GTGACTTCAG ACATTAGGTT CAGACATTAG TTAAAAATAG	1320
50	AAAGAGGAAT AAAGACATOT YTTOTOTOTA GAAAAGATAA CACCECAAST AATAATOOTT	1380
30	CCCACTITCA TIGAGATCAG CTIGICIGAT AACCTGATAI GAGTGACA ATGATAAACA	1440
,	TGATAATAGT GGTACTTTTG TAATTTTGCT GGTGCATTTA AGAAGATAGT AAAKGATGAG	1500
55	TYCAYCTTTT CTYCGAACAT YCCTATYCCT AGATGLAGTT TACCTCAAAT TGGGAATTAT	1560
	AACTGTCCTA ATTTTTGTTG TGTACCCTGA TGCCCCTTTT GCTTTAATAC CCACAGTGTA	1620
60	ACAATTAAAT ATCACACTAT GACATAIGAI TIAAGIAGGA TATTITAAAG ATAAATTTTA	1680
55		

	GGGGTAAATG	TTTACTICAA	AATGACTCCA	TATTTCAAAT	ATCTGTTTAG	ACTGTGAAGG	1740
	ССАААТААТТ	TTTAAGAAAA	CATTTGAAGA	GTAGTGTGTT	TGCATTIGIG	AATAATCTTA	1800
5	CTCACAGCAA	GTAAACGTAA	TAAAAGCCAA	CATTTAAGCC	АААААААА	ААААА	1856

10 (2) INFORMATION FOR SEQ ID NO: 53:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1558 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

#### (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:

20	TGGGTATCCA	TTCCTGNAAT	TACTTTACTT	AGGATAATGG	CCTCCAGCTC	CGTCCAAGTT	60
	GCTGCAAAAG	GTATTATTTC	GTTCCTTTTT	GTGGCTGAGT	AGTATTCCAT	GGTGTATATA	120
25	TACCACATTT	TCTTTATCCA	CTCATTGCTT	GATGGGCAGT	TAGGTTGGTT	CCACATCTTT	180
23	GCAATTGTGA	GTTGTGCTGC	TCCAGATATC	ATCTTTAACT	CCTTTGCCTŢ	CTCCACATAC	240
	ATTTCCAAGT	CCTGTTCATT	CTACCTCCAA	AATGTATCTT	GTATCCATTC	ATCTCTCTCC	300
30	ATCTTCAATC	TATTTCAATG	CCCCATCATC	TCTTGCATGG	AGGAGTGTAA	TAATTGGCTA	360
٠	ACTGGCCTGT	TCTTACATTT	таааатсааа	AGATGTGACA	GGTGAAATGC	CTATTTCAGT	420
35	GTCCATTGAT	GGTTCTGCTT	ACACACCACC	TGGCTGCCTG	GTGTCGCAGT	GGCAGAGTTG	480
33	AGCAGTGTGA	AAAAGACTGC	TTGGCCCTTT	ACAGGGAAAG	CAGGTCCACT	GTGGCCTGTG	540
	AGGACGAGAG	CTCTGGGCAG	GCTCGGACAC	TGGCAGACCC	TGGTCCTGGC	TGGCCAAGGC	600
40	AGCAGGGTAT	GTGTTTCGGG	TCACTCACAG	GGCTCAGCAC	CACTCCTCAT	GGCTTCCTTA	660
	CTGTTTCGGC	AGAGGCTGAC	CCGCGGCTGA	TTGAGTCCCT	CTCCCAGATG	CTGTCCATGG	720
45	GCTTCTCTGA	TGAAGGCGGC	TGGCTCACCA	GGCTCCTGCA	GACCAAGAAC	TATGACATCG	780
43	GAGCGGCTCT	GGACACCATC	CAGTATTCAA	AGCATCCCC	GCCGTTGTGA	CCACTTTTGC	840
•	CCACCTCTTC	TGCGTGCCCC	TCTTCTGTCT	CATAGITGTG	TTAAGCTTGC	GTAGAATTGC	900
50	AGGTCTCTGT	ACGGGCCAGT	TTCTCTGCCT	TCTTCCAGGA	TCAGGGGTTA	GGTGCAAGA	960
	AGCCATTTAG	GGCAGCAAAA	CAAGTGACAT	GAAGGGAGGG	TCCCTGTGTG	TGTGTGTGCT	1020
e-e-	GATGTTTCCT	GGGTGCCCTG	GCTCCTTGCA	GCAGGGCTGG	GCCTGCGAGA	CCCAAGGCTC	1080
55	ACTGCAGCGC	GCTCCTGACC	CCTCCCTGCA	GGGGCTACGT	TAGCAGCCCA	GCACATAGCT	1140
	TGCCTAATGG	CTTTCACTTT	CTCTTTTGTT	TTAAATGACT	CATAGGTCCC	TGACATTTAG	1200
60	TTGATTATTT	TCTGCTACAG	ACCTGGTACA	CTCTGATTTT	AGATAAAGTA	AGCCTAGGTG	1260

	TTGTCAGCAG GCAGGCTGGG GAGGCCAGTG TTGTGGGCTT CCTGCTGGGA CTGAGAAGGC	1320
5	TCACGAAGGG CATCCGCAAT GTTGGTTTCA CTGAGAGCTG CCTCCTGGTC TCTTCACCAC	1380
3	TGTAGTTCTC TCATTTCCAA ACCATCAGCT GCTTTTAAAA TAAGATCTCT TTGTAGCCAT	1440
	CCTGTTAAAT TTGTAAACAA TCTAATTAAA TGGCATCAGC ACTTTAACCA AAAAAAAAA	1500
10	AAAAAAAAA AAANAAAAA AAAAGGGGC CGCTCTAGAG GTCCAAGTTA NGACGNGG	1558
15	(2) INFORMATION FOR SEQ ID NO: 54:	•
20	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 948 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:	
25	TAAAAATCAT GCTCTGTACC ATCCTCACCG TAGTCATCAT CATCGCCGCG CAGACCACGA	60
•	GAACTACTGG GATCCCTAAA AACGCCCCTG GTCCGGCCCC ACTCTGCGCC CCTCGATCTC	120
30	CCAGGCTCTT TCTGCAGWCA TACCGCGGAC CCAATGGGCG CCCTGCACAC CCGTTTCTGG	180
	GGCCGTCAGA CTTGGATACA TCGTAAACTC CGCCTCCACG GAACGTCTCG CCTKGCGAGC	240
	AAGMTCGGAA TCCAGTTCCT CAGGAACCCC TCCAAAACCC ACACCCCCAG GGACGCCGCT	300
35	TTCCGGGATC CCGGSCAAAC GCCGGACCCT CAGTCGCTCC AGGCCCCCTC ACCCTCAAAG	360
	TGTAGCGCCC CCAACCGAGC AACCTCGGTT TGGTCCCTAA AACCCCGCCT CCTCTATAAG	420
40	CACCGCCCCA GCTCTGACAA AACCCCGCCT CCAGGTCGGC AGGCTCCGCT TCTTTTCTTC	480
	TCCGCGGGGT GATTCAGTCC AGTGATTGGG TTTGTGGCTC CAGGCCTCGC CCACAGACGG	540
	ACAGACCCCT CCCTTTCTTC CGGCAAAAGG ACCGAGCCCT GGGGTAGTAA GGSCCCCACA	. 600
45	CTCCTGTTTT TTGCAAGTAC ATTTTTGTCC YTCCTCCACC CAGGTATCTG CCTATTTTCT	660
	TGCTAATCCC AGAACCTTTC CTTTTGCTTT TTTTAAGGAC ATTTGGGAAG TTCCTGGTGT	720
50	AGGACCCTTC TCCCTGGGAT AAGAAACCTG CCTGTAAACG CTCTGTAAAT ACTCCCTTCC	780
- •	ACCCATCCCA GCCCCTGGGC AGCCGGGCAG AAGGGAATCC AGGCTATGGA CCTCCCAAGT	840
	CCCCGCTCCC CGCTCCCCTC GGCGGCCCCG CCTTGTTCTG ATCTGTGTGT GAGTGTGTGT	900

GAACTICIGA AAGACAATAT TAAAGAGACT TAGTIGAAAA AAAAAAAA

5	(A) LENGTH: 990 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:	
10	GGGGAACTGC AGTGACAGCA GGAGTAAGAG TGGGAGGCAG GACAGAGCTG GGACACAGGT	60
	ATGGAGAGGG GGTTCAGCGA GCCTAGAGAG GGCAGACTAT CAGGGTGCCG GCGGTGAGAA	120
15	TCCAGGGAGA GGAGCGGAAA CAGAAGAGGG GCAGAAGACC GGGGCACTTG TGGGTTGCAG	180
15	AGCCCCTCAG CCATGTTGGG AGCCAAGCCA CACTGGCTAC CAGGTCCCCT ACACAGTCCC	240
	GGGCTGCCCT TGGTTCTGGT GCTTCTGGCC CTGGGGGCCG GGTCGGCCCA GGAGGGGTCA	300
20	GAGCCCGTCC TGCTGGAGGG GGAGTGCCTG GTGGTCTGTG AGCCTGGCCG AGCTGCTGCA	360
	GGGGGGCCCG GGGGAGCAGC CCTGGGAGAG GCACCCCCTG GGCGAGTGGC ATTTGYTGCG	420
25	GTCCGAAGCC ACCACCATGA GCCAGCAGGG GAAACCGGCA ATGGCACCAG TGGGGCCATC	480
	TACTTCGACC AGGTCCTGGT GAACGAGGGC GGTGGCTTTG ACCGGGCCTC TGGCTCCTTC	540
	GTAGCCCCTG TCCGGGGTGT CTACAGCTTC CGGTTCCATG TGGTGAAGGT GTACAACCGC	60 <u>0</u>
30	CAAACTGTCC AGGTGAGCCT GATGCTGAAC ACGTGGCCTG TCATCTCAGC CTTTGCCAAT	660
	GATCCTGACG TGACCCGGGA GGCAGCCACC AGCTCTGTGC TACTGCCCTT GGACCCTGGG	720
35	GACCGAGTGT CTCTGCGCCT GCGTCGGGGG NAATCTACTG GGTGGTTGGA AATACTCAAG	780
	TTTCTCTGGC TTCCTCATCT TCCCTCTCTG AAGGACCCAA GTCTTTCAAG CACAAGAATC	840
	CAGCCCCTGA CAACTTTCTT CTGCCCTCTC TTGCCCCCANA AACAGCANAA GCAGGANANA	900
40	NACTOCCTCT GGCTCCTATC CCACCTCTTT GCATGGGAAC CTGTGCCAAA CACCCAAGTT	960
	TAAGAAAAAA ATAAAACTGT GGCATCTCCA	990
45		•
	(2) INFORMATION FOR SEQ ID NO: 56:	
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1603 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:	
	GGTCGACCCA CGCGTCCGGC CCGCCGGCTC CGGAGCGGCT CTGCCTTCCC GAGCGCGGGA	60
60	CCGCGCCCTG GGGGAGGAGG GCGAACGACG CGGCGATGGC TCCGCGGGCA CTCCCGGGGT	120

•	CCGCCGTCCT	AGCCGCTGCT	GTCTTCGTGG	GAGGCGCCGT	GAGTTCGCCG	CIGGIGGCIC	180
	CGGACAATGG	GAGCAGCCGC	ACATTGCACT	CCAGAACAGA	GACGACCCCG	TCGCCCAGCA	240
5	ACGATACTGG	GAATGGACAC	CCAGAATATA	TTGCATACGC	GCTTGTCCCT	GTGTTCTTTA	300
	TCATGGGTCT	CTTTGGCGTC	CTCATTINGC	CAMCTNGCTT	NAAGAAGAAA	GCTATCGTT	360
10	GTACAACAGA	AGCAGAGCAA	GATATCGAAG	AAGAAAAAGG	TIGAAAAGWI	AGRATTGAAT	420
10	GACAGTGTGA	ATGAAAACAG	TGACACTGTT	GGGCAAATCG	TCCACTACAT	CATGAAAAAT	480
	GAAGCGAATG	CTGATGTYTT	AAAGGCGATG	GTAGCAGATA	ACAGCCTGTA	TGATCCTGAA	540
15	AGCCCCGTGA	CCCCAGCAC	ACCAGGGAGC	CCGCCAGTGA	GTCCTGGGCT	TTGTCACCAG	600
	GGGGGACGCC	AGGGAAGCAC	GTCTGTGGCC	ATCATCTGCA	TACGGTGGGC	GGTGTWGTCG	660
20	AGAGGGATGT	GTGTCATCGG	TGTAGGCACA	AGCGGTGGCA	CTTTATAAAG	CCCACTAACA	720
	AGTCCAGAGA	GAGCAGACCA	CGGCGCCAAG	GCGAGGTCAC	GGTCCTTTCT	GTTGGCAGAT	780
	TTAGAGTNAC	AAAAGTGGAG	CACAAGTCAA	ACCAGAAGGA	ACGGAGAAGC	CTGATGTCTG	840
25	TTAGTGGGGC	TGAAACCGTC	AATGGGGAGG	TGCCGGCAAC	ACCTGTGAAG	AGAGAACGCA	900
	GTGGCACAGA	GTAGCAGGTG	AGCCGTGGTT	TTGGTGACAT	TGGGGGCAGA	GTGGTGCAGG	960
30	GTGAGGAGAA	GGTACTTGGA	GCCTCCCAGG	TGCTGTGGCA	GCATAGGAAT	GGTATTTGAC	1020
	AGGGAAGTGG	GAGAGCTTTC	CTTGACCCAG	GAAGACTGAG	GGGGACTGAA	CATGATTACT	1080
	TGTCTGCCTA	GAGCTTCTTG	TAAAGAAGTC	ACAAACTTAG	TGCCTCCAGG	GGCTTGGCTG	1140
35	TGTGATAATG	AGGATAGAGG	ATTACTTGTG	AGGCAATGTG	GCATGGTGGG	GATTGTGGCA	1200
	AACTAGAATT	CACATCACCC	ACCATATAGG	GCTTGCATTA	CCACGAGGCA	GAAAGCACCT	1260
40	AGTGTTGCTG	CATCTTCTTA	CGCAAAAAAG	ACAAAATCCA	GACTTCTAAA	ATGTAAAATC	1320
	ACTGATTTTC	GATATTGGCA	GCTTACTTTT	TTTTTTTAAA	CAACCATGCA	GGCCAAATGA	1380
	CTTGTAATCT	TGTCACCATT	TTTAGGTAAA	CTGTGACTTG	AAAAAGTCTG	GAGCAAACAA	1440
45	ACCAATGCTT	TTTCCTTTTA	TTCTGTTGGR	AACCAGTTTT	CTTTGTGTCA	CAGTTYTGAA	1500
•	ACCTCAATAC	GAATATTTCT	CTTCCCACCA	AATATTTTGA	GGCAATTGAA	AAGCCACAGT	1560
50	GATTTATTTC	TTGATTTGGC	AATTTTAATT	TTGCAAGACA	ATT		1603

(2) INFORMATION FOR SEQ ID NO: 57:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1052 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5	7				•									į		٠			j	j	j	j	į	į	į	į	į				į	,	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	į	,	,	,	5		2									:		)	2	٥	(	Į	J	1			J	)	I		I	]			,	2	2	Ç	)	3	E	1	5	3	ŝ	٤	S	:	:					:	ľ	Į	V	١	1	)	)		C	(		C	Į	Į	J	J		'	•	•	ľ	l	l	)	•	•	•		Ŀ	)	1		•		I
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5	TACAGCTCAG	GATGCCTGTA	ACATTGTCAT	CTCTGGGCTT	CTGGGTCCTG	CTTAGCCTGC	60
J	TTTTTCCCTG	GAGGACTGAC	CAGGGATGCG	GCCCAGCAAC	ATGTTACTAA	ATCATACTCT	120
	CCTCCCTACC	TTTCCCAGAC	CTCTCACTCC	TGCCTGGTGT	TCCÁACCCGT	TCTGTGGCCA	180
10	GAGTATACAT	TTTGGAACCT	CTTCGAGGCC	ATCCTGCAGT	TCCAGATGAA	CCATAGCGTG	240
	CTTCAGCAGN	AAGGCCCGAG	ACATGTATGC	AGAGGAGCGG	AAGAGGCAGC	AGCTGGAGAG	300
· 15	GGACCAGGCT	ACAGTGACAG	AGCAGCTGCT	GCGAGAGGGG	CTCCAAGCCA	GTGGGGACGC	360
13	CCAGCTCCGA	AGGACACGCT	TGCACAAACT	CTCGGCCAGA	CGGGAAGAGC	GAGTCCAAGG	420
	CTTCCTGCAG	GCCTTGGAAC	TCAAGCGAGC	TGACTGGCTG	GCCCGTCTGG	GCACTGCATC	480
20	AGCCTGAATG	AGGCTGGCCA	CCTGCCACTT	TGCCCTGCCC	TCTGCCTCCA	GGGCTCCMCT	540
	муссттсстт	TTCTTGGTGA	AAGGCACCTC	CTTTCCTGAT	AATGAATGGT	GTTCCCTTTG	600
25	CTTGGCTGGG	GAGCCCCCA	GGCCAGGTTT	GCTGGCCATA	GATACCTTTG	GGCTGCCTGR	660
23	GACAGGCTCC	TGAGGAGGAT	TGAGGGTGAA	AGTCTCCCAC	GAGTACACTA	AACCTAGGTC	720
	TGGTCACCAA	TAGGGTTTGG	AGAGCAAAGG	GCCACAACTC	ATCAGCTGCC	TGTCTCTTAG	780
30	ATGCACTTTC	TTTTTCCACC	AGCACATCCT	TCAACACACA	GAATTTCAGG	GAAGAGTTCT	840
	CCCCAAAACC	CTAGCTCTTT	ACCCTTCCAT	TTTAGCCTTC	CACCCAGCTT	CCACAAAAGA	900
35	TTTGGCTCTA	CCTTGGATCT	GCTAGTAAAT	AACTAATAGG	CAGGCAGTTA	TTTGGGTAAG	960
	· GAAAAAAGGG	GTGGGAGAGA	CAGAAAATTT	GCCCACTGCT	GCTCCTCCCC	TTGGSTYTCC	1020
	ACCTGGGATT	TGCTATTGAA	TCTCTACCCT	NN			1052

(2) INFORMATION FOR SEQ ID NO: 58:

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45 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 814 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:

•	ACTGCAACCT GGGAGGCAGA GGTTGCAGTG AGTCGAGATG GTGCCATTGC TCTCGTTTGG	30
	GCAACAAGAG TGAAACTCTT GTCTCAAAAA AAAAAAAAAA	36
5	GTCATTACTG GTGGGATCTG GTCACACAAG ATAGCATTAA ACGTGACATG GCACATAAAA	42
	TIGGITAAAA AATITIGITT TITAATTACG TAATGTAAAA GCCCAACAAA CACTITATGC	48
10	AAGATTGGAA TGTATCTTCA AATTCAGATT TAATAAACAT GTAAAGATCC TCTGTATATA	540
10	AAAGTTGTAT TTAATCCCTT GTGCCCCAAG AATGCTATAA AAGATCCCAA GAATGTTATC	600
	TATGAAAAGA TAGCAATAGG GAATGGTGAA CAAATAATTT AATTTGCCAA TTCTAAAAAA	660
15	CATGGACTTA AACCCCATGA AAACTTGGTT CCATAGTTTT AACTGTTTTA TGGTTCCAAT	720
	ACAAAACCAG AGTGGTTTAC ATTCCACAAT NACCAAATTT GCATCCAATN TTGGGGTAAT	780
20	TTTNGGTATT TGCCATGGGA TACTATTCAT TTTT	814
20		
25	(2) INFORMATION FOR SEQ ID NO: 59:	
23	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1215 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
30	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:	
35	AGAGGAAGTC TTTTGCCAAG CCTGTTCTCT GGACTAACGC CATCCAGGCT GGGAGGGGAA	60
	GAGTGCTCTG CTACACTCGT CCCCCTCCTG CCTCATCTTC CTTCTCAGCC TTGGTTCCTG	120
	ATGGGAACAG AATGGAGGGC CTGAGAACAT ACTTTCTAAA TGCCTTTGAC CCAGGAACCG	180
40	ATTATCTATA TTTGTTCCCA TTTTCCTTCA CCGTGACATT CCAGCATTGT CTGACTGTGA	240
	GGTGGGCCTT TGAGAGCCTC CAGGTTCCTC AAAACAGGCC TGAGCGATGG GCATCACACC	300
45	CTCTGCCTAC CCACRTGCCT GCTTACCTGC CAGATAACCA AGTGNAGATG TCTGCGAGTG	360
	GCTAGTTTTC ACATTCTTAC TAGTGTTTGG YTCACCTTTG GGCAAAGGCC CCCTCTAGGC	420
	CTTGCCCCAC CTCCATCAAA CGCAGACACT GTAGTCAGAC CTCAGYAATA TAGGAGGCAA	480
50	TAATCTTTTA ACAGTGTTTT GCAAACAAAC AAAAAGAGAA AAATCCCAGC CAGGGGAACT	540
	CGCCACCTGC CCACGCTAGT TCCATCCACG CTCAAGACCC GCCCTTAGAC CAGGCAGGCA	600
55	AAGGCCCCCA TCACACTCGG CCACTAGTGG GGTCCTGAGG CCAAGAAAGA AACCAGACCC	660
55	TGTATGACAA GTTGGGKTCT TTCCAGAACA CGACAGAAAC AGGGGGGGCC CCTTGTTAAT	720
	GCCACTCCAT ACTCCAGAAG CATTATTCCT TATTTGGGAC AGCCAAGGGC AGATTCACAG	780
60	GTTATTGTAG GAATAAAGAC TAGTTTACAA AGGARAAAGA GSCCCTGGAC TTCCCMAGGA	840

•	AAGGTCAGGT TAGGGCTCCT GTACCCATTC TGTTCCACCA CTGTTTGATC TCTCTGGCCT	900
5	CCCACCAGGA ATGCCGTTTC CTTTTTATGG ATCTGTTGGG AACCAGAGAG AATCAACAGA	960
5	TCAATGACAT AGGATCCGAA GTGCAATGAT AGTCACTTCT AGTTTGGCAT TTCACAAACT	1020
	CTGNACAGCA AGGTATTGGT AGGTTACTCA ATTTCAAAAG GGCCCCATGG CCAAATATGT	1080
10	TTAGGAACCG CTGTTTGNAT TTCTTTTTTT GGAGACGCAT TGTATATAAT ATATGTCAAA	1140
	GGCTTTCGGA ATTCCTGCAG GAAAGAAATC AGCTTTGTTA AATCCNAAAA AAAAAAAAAAA	1200
15	AAAAAAATAG ACTCG	1215
13		
20	(2) INFORMATION FOR SEQ ID NO: 60:  (i) SEQUENCE CHARACTERISTICS:	
25	<ul><li>(A) LENGTH: 478 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li><li>(D) TOPOLOGY: linear</li></ul>	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:	
30	ATTTCTTATG ACATGGGGGT TTGAATTGGT TGGCAAATGT TTAATTTTAA TATCCATAAT	60
	CAGTGAGGTC CTGCTGGCTG TAATCATTAA TTGTGAAATC TAAGGAGCTT AGTTCATGGC	120
	TCTAGAATTT CACAGAAAAR TGYGMTATGA TACGAGCATT AAGTTTATTT CTTCTGATCT	180
35	TTGATGCAGC TTTGTTCAGT TTATCTGTTT TTGTATTTAT TGGTCATCTA CTTCCCATGC	240
	CAAAAGGGAC TGGTCTACAT AGCTGCGCTA AACACCTGAT CAAATCACTA AAAGAAAATG	300
40	TGTTACCTCT AATGAATTAT CCTGATTGTA AGTTAAAAAT CAATATTTCC CCGTAGTGAG	360
	GTTTGCTTTT TAAAAAGAAK KCTTAAAAAA AAAAAAAAAA AAACGAGTTN AAGAAAAGGA	420
45	AGCAAGCTCA GGTAAGGTGC ACACATTGGG CTAAGGAAGC TAGAGCCTGT GGAGANGC	478
	(2) INFORMATION FOR SEQ ID NO: 61:	
50	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 618 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li></ul>	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:	
60	TATGACCTTG ATAACCCCAA GTTNGAAATT AACCTTCANT AAAGGGAACA AAAGCTGGAG	60

WO 98/54963 PCT/US98/11422

GCTACTAGGT AAGCCTTCTG GGACTTTCAG ATATTTTGGG GAAGATTGAT TTTTGTTCTT  ACATGCTGTG GACCCTTGGC CATCAAATGG TATGGGGAAG CTCATCCGTC TGTCTGTGAT  GGTCATGTCA GTCAGGCGTC TTTTTAGTAT TTACTGGGTAG CTCAGTACTG TGCCAGATGC  TGTCGGGAGC CGTGGTGGTA TGGAGGAGGA GTGCTCCAGA GGACTCTGCT GTGTGGCAGG  CCAGCATAAA CAAGCCAAGG GGAAAAGGCA GGCATGGAAT AAAGGGGGAG AATACCAGTG  TGTGACTTAC TGCTGACTGT GTGGATTAGC CTATCAGCAG TAATCAAGCA GGGCGGAGGG  CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG  TGAGACCACA AGGCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  (2) INFORMATION FOR SEQ ID NO: 62:  (A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  35. TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTACAAGGAG ACTACGATGC CTGCCTTGGT CACCCTTCTC CTGCTCTTC CTGAGCTGAG	180
GGTCATGTCA GTCAGGCGTC TTTTTAGTAT TTACTGGGTG CTCAGTACTG TGCCAGATGC  TGTCGGGAGC CGTGGTGGTA TGGAGGAGGA GTGCTCCAGA GGACTCTGCT GTGTGGCAGG  10 CCAGCATAAA CAAGCCAAGG GGAAAAGGCA GGCATGGAAT AAAGGGGAG AATACCAGTG  TGTGACTTAC TGCTGACTGT GTGGATTAGC CTATCAGCAG TAATCAAGCA GGGCGGAGGG  CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG  TAGGACCNCA AGGCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  (2) INFORMATION FOR SEQ ID NO: 62:  (A) LENSTH: 751 base pairs (B) TYFE: nucleic acid (C) STRANBEMESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  35 TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTC ACCCCTGGAC  CTAGTGGACTC ACTTCTAACA ANGAGAATAC AGCACANAT GTAAAATTC ACCCCTGGAC  CTACAAGGGA ACTACGATGC CTGCCTTGGT CACCCTTCTC CTGCTCTTTC CATTGCTCCC  TCTGATGGAA GCCACTTGCC ATGTGATGAG GTGCCCTATG GAGAGGCCCA CGTGACAGGG  45 TATTGTAAAA AGCCTCTGAC CAATAGCCAT CTAGAAACGG AGGCCCAGTC CAGCAGCCTC  TGAGATGAAT CCTGCCCAAC CAATAGCCAT CTAGAAACGG AGGCCCAGTC CAGCAGCCTC  TGAGATGAAT CCTGCCCAACC TGAGCTTGGAG GACCCAAGCCA GTGAACCCAAA  TGATCACAGC CACCACCAAC ACCTTCCACTG CCTGGTGAGA GGCCCAAGCCA GTGAACCCAA	
GGTCATGTCA GTCAGGCGTC TTTTTAGTAT TTACTGGGTG CTCAGTACTG TGCCAGATGC  TGTCGGGAGC CGTGGTGGTA TGGAGGAGGA GTGCTCCAGA GGACTCTGCT GTGTGGCAGG  10 CCAGCATAAA CAAGCCAAGG GGAAAAGCCA GGCATGGAAT AAAGGGGGAG AATACCAGTG  TGTGACTTAC TGCTGACTGT GTGGATTAGC CTATCAGCAG TAATCAAGCA GGGCGGAGGG  CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG  15  TAGGACCACA AGGCTTCTIN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  20  (2) INFORMATION FOR SEQ ID NO: 62:  (A) LENGTH: 751 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  35, TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	240
10 CCAGCATAAA CAAGCCAAGG GGAAAAGGCA GGCATGGAAT AAAGGGGGAG AATACCAGTG TGTGACTTAC TGCTGACTGT GTGGATTAGC CTATCAGCAG TAATCAAGCA GGGCGGAGGG CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG 15 TAGGACCNCA AGGCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC AGTTTTGGGA AGCAAGGG 20 (2) INFORMATION FOR SEQ ID NO: 62:  (i) SEQUENCE CHARACTERISTICS:	300
TGTGACTTAC TGCTGACTGT GTGGATTAGC CTATCAGCAG TAATCAAGCA GGGCGGAGGG  CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG  TAGGACCNCA AGCCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  (2) INFORMATION FOR SEQ ID NO: 62:  (i) SEQUENCE CHARACTERISTICS: (ii) LENGTH: 751 base pairs (iii) TYPE: nucleic acid (ii) STRANDENNESS: double (iii) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  ATGGCCCTGA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	360
CATTATCTTT GAGCCAGAAG AGTGAGCACT GGSCCGAGGG TGGAGCATCA AGAGGGGGTG  TAGGACCNCA AGGCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  (2) INFORMATION FOR SEQ ID NO: 62:  (a) LENGTH: 751 base pairs (b) TYPE: nucleic acid (c) STRANDEDNESS: double (d) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	420
TAGGACCNCA AGGCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  (2) INFORMATION FOR SEQ ID NO: 62:  (3) SEQUENCE CHARACTERISTICS: (A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	480
TAGGACCNCA AGGCTTCTTN CNGGGGAGAC AACGTCAATA AGCNGTCAGT AGTCACCGAC  AGTTTTGGGA AGCAAGGG  (2) INFORMATION FOR SEQ ID NO: 62:  (3) SEQUENCE CHARACTERISTICS: (A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	540
(2) INFORMATION FOR SEQ ID NO: 62:  (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TOGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	600
(2) INFORMATION FOR SEQ ID NO: 62:  (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	618
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 751 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 751 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	
(A) LENGTH: 751 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear  (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	
(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	•
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	
(xi) SEQUENCE DESCRIPTION: SEQ 1D NO: 62:  TCGACCCACG CGTCCGAGGA GCTGGACTTC TGAGACAGCC ATTCTCCTTG CATAGCACTG  35, TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA  ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	
TCTGCTGCTA CAGCTCATAG AAGTCAACAA TTTTCTTCAA CACTGGTAGG CAGCCTCTAA ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	
ATGGCCCTGA TCACCCTCAC CTCCTGCCAT TCACACCNNT GTAAAATTCC ACCCTGGAC  CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	60
CTAGTGACTC ACTTCTAACA ANGAGAATAC AGCAAAAGTA ACATCGCTTC TGAGGTGAGG	120
CTACAAGGAG ACTACGATGC CTGCCTTGGT CACCCTTCTC CTGCTCTTTC CATTGCTCCC  TCTGATGGAA GCCAGTTGCC ATGTGATGAG GTGCCCTATG GAGAGGCCCA CGTGACAAGG  TATTGTAAAA AGCCTCTGAC CAATAGCCAT CTAGAAACGG AGGCCCAGTC CAGCAGCCTC  TGAGATGAAT CCTGCCAACC TGAGCTTGGA GACAGATTCT CTCCCTATCC TGCCTTGGGA  TGATCACAGC CACCACCAAC ACCTTCACTG CCTGGTGAGA GGCCAAGCCA GTGAACCCAA	180
CTACAAGGAG ACTACGATGC CTGCCTTGGT CACCCTTCTC CTGCTCTTTC CATTGCTCCC  TCTGATGGAA GCCAGTTGCC ATGTGATGAG GTGCCCTATG GAGAGGCCCA CGTGACAAGG  TATTGTAAAA AGCCTCTGAC CAATAGCCAT CTAGAAACGG AGGCCCAGTC CAGCAGCCTC  TGAGATGAAT CCTGCCAACC TGAGCTTGGA GACAGATTCT CTCCCTATCC TGCCTTGGGA  TGATCACAGC CACCACCAAC ACCTTCACTG CCTGGTGAGA GGCCAAGCCA GTGAACCCAA	240
TATTGTAAAA AGCCTCTGAC CAATAGCCAT CTAGAAACGG AGGCCCAGTC CAGCAGCCTC TGAGATGAAT CCTGCCAACC TGAGCTTGGA GACAGATTCT CTCCCTATCC TGCCTTGGGA TGATCACAGC CACCACCAAC ACCTTCACTG CCTGGTGAGA GGCCAAGCCA GTGAACCCAA	300
TGATCACAGC CACCACCAAC ACCTTCACTG CCTGGTGAGA GGCCAAGCCA GTGAACCCAA	360
TGATCACAGC CACCACCAAC ACCTTCACTG CCTGGTGAGA GGCCAAGCCA GTGAACCCAA	420
	480
	540
50 GGTAAACTGG ACAGAATCCT GACCCACAGA AACTGAGATA ATGTTTGTTA TTTTAAGCTG	600
CTCAGTTTGT TACAGAGCAA TAGATAACTA ACTCAAACAC CATAAAATTC TAATATTTTA	660
55 TTCTATCACA CAAACCAGGT AATACCAAGT AAATGCCATT ACTATACACA TATTTTTGTA	720
ACACAATTAC ATGTGATTTT TTAAGAAGGC T	

	•	
•	(2) DEFORMATION FOR SEQ ID NO: 63:	
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 780 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPCLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63:	-
	CNGNCLETCA CNGTCCCCGA TTCCCGGGTC GACCCACGCG TCCGGGTTGG CAACTCCTGA	60
15	GGCCTGCATG GGTGACTTCA CATTTTCCTA CCTCTCCTTC TAATCTCTTC TAGAGCACCT	120
13	GCTATCCCCA ACTICTAGAC CIGCTCCAAA CTAGTGACTA GGATAGAATT TGATCCCCTA	180
	ACTCACTGTC TGCGGTGCTC ATTGCTGCTA ACAGCATTGC CTGTGCTCTC CTCTCAGGGG	240
20	CAGCATGCTA ACGGGGGGAC GTCCTAATCC AACTGGGAGA AGCCTCAGTG GTGGAATTCC	300
	AGCCACTGTG ACTGTCAAGC TGGCAAGGGC CAGGATTGGG GGAATGGAGC TGGGGCTTAG	360
25	CTGGGAGGTG GTCTGAAGCA GACAGGGAAT GGGAGAGGAG GATGGGAAGT AGACAGTGGC	420
23	TEGTATEGOT CTEAGGCTCC CTEGGGGCCTG CTCAAGCTCC TCCTGCTCCT TGCTGTTTTC	480
	TGATGATTTG GGGCCTTGGG ASTCCCTTTG TCCTCATCTG AGACTGAAAT GTGGGGATCC	540
30	AGGATGGCCT TECTTECTCT TACCETTECT CECTEAGECT GEAACCTCTA TECTGGAACC	600
	TGTCCTCCCT TTCTCCCCAA CTATGCATCT GTTGTCTGCT CCTCTGCAAA GGCCAGCCAG	660
35	CTTGGGAGCA GCAGAGAAT AAACAGCATT TCTGATGCCA AAAAAAAAAA	720
33	GCGGCCGAAA GCTTATTNCC CTTTAAGTAA GGGGTTAATT TTTAGCTTGG GCACTNGGCC	780
40	(2) DIFOFMATION FOR SEQ ID NO: 64:	
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 588 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 64:	
U	TTCCGAATTA ATCGACTCAC TATAGGAAWT GCCGTCGCCA TGACCCGCGG TAACCAGCGT	60
	GAGCTCGCCC GCCAGAAGAA TATGAAAAAG CAGAGCGACT CGGTTAAGGG AAAGCGCCGA	120
55	GATGADGGGC TTTCTGCTGC CGCCCGCAAG CAGAGGGACT CGGAGATCAT GCAGCAGAAG	180
	CAGAAAAGG CAAACGAGAA GAAGGAGGAA CCCAAGTAGC TTTGTGGCTT CGTGTCCAAC	240
	CCTCTTGCCC TTCGCCTGTG TSCCTGGAGC CAGTCCCACC ACGCTCGCGT TTCCTCCTGT	300

	AGTICCTCACA GGTCCCAGCA CCGATGGCAT TCCCTTTGCC CTGAGTCTGC AGCGGGTCCC	360
	TITTGTGCTT CCTTCCCCTC AGGTAGCCTC TCTCCCCCTG GGCCACTCCC GGGGGTGAGG	420
5	GGGTTACCCC TTCCCAGTGT TTTTTATTCC TGTGGGGCTC ACCCCAAAGT ATTAAAAGTA	480
	ССТТТСТААТ ТССААЛАЛАА АЛЛАЛАЛАА АЛЛАЛАЛАА АЛЛАЛАЛАА АЛЛАЛАЛАА	540
10	AAAAAAAAAA AAAAAAAAA AAAANNCGGG GGGGGCCCC CCCCCCCC	. 588
10		
	(2) INFORMATION FOR SEQ ID NO: 65:	
15		
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 774 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:	
25	TTTAAAGATG AAGAAATGAC AAGGGAGGGA GATGAGATGG AAAGGTGTTT GGAAGAGATA	60
23	AGGGGTCTRA GAAAGAAATT TAGGGCTCTG CATTCTAACC ATAGGCATTC TCGGGACCGT	120
	CCTTATCCCA TITAATTAAT TICTCTGACA ATTCAATTAT TITCTGTTAT TAATGTTGCC	180
30	ACTGCTTTCT GTTTGTCTGC ACTTTCTTGA TAAATATTTG CTATCGTTTT ACTCCAGTCA	240
	TTCGATGTTG CTGAGATTTA CATATGACTC TTGTCAACAT CTCATCTTTT GACCCAATCT	300
35	TATTCATTTA ATAAGAGGTC TCATTCATTT GCATGGAAAA ATGCTCATTG TATATTGCAA	360
<i>J J</i>	AGTGAAAATA ACGAGTTGCA AAACAGTGTA TACATATATG TGTGTATATA TGTACACTTT	420
	ATTTGTACAT TTCTATGTGA CATAATGCAA AGGAAAGTGT CTGATTTTAT TATACACCAA	480
40	AGGTTAACAG TGAATCTCTG TGTGATCTCT TTTTTTTCT TTTTGCCTAT CTGCATCTTC	540
	TCACTTGCCA AAAAATGAAT ATATGTTTAT GTGTGTATAT TACTTGTGTC ACAAAAAACC	600
45	CTAAAGTAGA CAGTAAAAGA ACTTGTCAAT CGCCTTTGGA AGGCAATGAA ACACTTAATA	660
-	AACTCTCAAT AACAGAAGCG TAAAAATGAA ATGTAAACCT CCAATTACCT CTGGATCTCT	720
	TAGCCAGAGT AATAAACTGG TAATTATTAC AGATAAAAAA AAAAAAAAAA	774
50		•
	(2) INFORMATION FOR SEQ ID NO: 66:	
55	(i) SEQUENCE CHARACTERISTICS:	

(A) LENGTH: 1866 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:

	•	(XI) SEQUENCE DESCRIPTION. SEQ ID No. 00.	
		ACCCACGCGT CCGGTCCTCT TCTTCAGCAC ATGCCAAAGC TGTTCCTCAC GGCCTGTGAG	60
	5	ACAAGAGCAT CTTGGATGTA GGACAATGGA AGAGTTAGAT GCCTTATTGG AGGAACTGGA	120
		ACGCTCCACC CTTCAGGACA GTGATGAATA TTCCAACCCA GCTCCTCTTC CCCTGGATCA	180
10	10	GCATTCCAGA AAGGAGACTA ACCTTGATGA GACTTCGGAG ATCCTTTCTA TTCAGGATAA	. 240
	10	CACAAGTCCC TTGCCGGCGC ANTCGTGTAT ACTACCAATA TCCAGGAGCT CAATGTCTAC	300
		AGTGAAGCCC AAGAGCCAAA GGAATCACCA CCACCTTCTA AAACGTCAGC AGCTGCTCAG	360
	15	TTGGATGAGC TCATGGCTCA CCTGACTGAG ATGCAGGCCA AGGTTGCAGT GAGAGCAGAT	420
		GCTGGCAAGA AGCACTTACC AGACAAGCAG GATCACAAGG CCTCCCTGGA CTCAATGCTT	480
	20	GGGGGTCTSG AGCAGGAATT GCAGGACCTT GGCATTGCCA CAGTGCCCAA GGGCCATTGT	540
	20	GCATCCTGCC AGAAACCGAT TGCTGGGAAG GTGATCCATG CTCTAGGGCA ATCATGGCAT	600
		CCTGAGCATT TTGTCTGTAC TCATTGCAAA GAAGAGATTG GCTCCAGTCC CTTCTTTGAG	660
	25	CGGAGTGGCT TGGNCTACTG CCCCAACGAC TACCACCAAC TTTTTTCTCC ACGCTGTGCT	720
		TACTGCGCTG CTCCCATCCT GGATAAAGTG CTGACAGCAA TGAACCAGAC CTGGCACCCA	780
	30	GAGCACTTCT TCTGCTCTCA CTGCGGAGAG GTGTTTGGTG CAGAAGGCTT TCATGAGAAG	840
	50	GACAAGAAGC CATATTGCCG AAAGGATTTC TTAGCCATGT TCTCACCCAA GTGTGGTGGC	900
		TGCAATCGCC CAGTGTTGGA AAACTACCTT TCAGCCATGG ACACTGTCTG GCACCCAGAG	960
	35	TECTITETIT GIGGGGACIG CITCACCAGI TITICIACTG GCTCCTICIT TGAACTGGAT	1020
		GGACGTCCAT TCTGTGAGCT CCATTACCAT CACCGCCGG GAACGCTCTG CCATGGGTGT	1080
	40	GGGCAGCCCA TCACTGGCCG TTGTATCAGT GCCATGGGGT ACAAGTTCCA TCCTGAGCAC	1140
	10	TTTGTGTGTG CTTTCTGCCT GACACAGTTG TCGAAGGGCA TTTTCAGGGA GCAGAATGAC	1200
		AAGACCTATT GTCAACCTTG CTTCAATAAG CTCTTCCCAC TGTAATGCCA ACTGATCCAT	1260
	45	AGCCTCTTCA GATTCCTTAT AAAATTTAAA CCAAGAGAGG AGAGGAAAGG GTAAATTTTC	1320
	į	TGTTACTGAC CTTCTGCTTA ATAGTCTTAT AGAAAAAGGA AAGGTGATGA GCAAATAAAG	1380
	50	GAACTTCTAG ACTTTACATG ACTAGGCTGA TAATCTTATT TTTTAGGCTT CTATACAGTT	1440
	50	AATTCTATAA ATTCTCTTC TCCCTCTCTT CTCCAATCAA GCACTTGGAG TTAGATCTAG	1500
		GTCCTTCTAT CTCGTCCCTC TACAGATGTA TTTTCCACTT GCATAATTCA TGCCAACACT	1560
	55	GGTTTTCTTA GGTTTCTCCA TTTTCACCTC TAGTGATGGC CCTACTCATA TCTTCTCTAA	1620
		TITGGTCCTG ATACTTGTTT CTTTTCACGT TTTCCCATTT CCCTGTGGCT CACTGTCTTA	1680
	- 60	CAATCACTGC TGTGGAATCA TGATACCACT TTTAGCTCTT TGCATCTTCC TTCAGTGTAT	1740
	00		

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TTTTGTTTTT	CAAGAGGAAG	TAGATTTTAA	CTGGACAACT	TTGAGTACTG	ACATCATTGA	1800
TAAATAAACT	GCCTTGTGGT	TTCAATAAAA	AAAAAAAA	АААААААА	АААААААА	1860
АААААА						1866

10 (2) INFORMATION FOR SEQ ID NO: 67:

### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1152 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:

			•				
20	CTCAAGGATG	TAAAGGCTCT	GCAGATTTCG	GGAGGCCTGT	CTCCCAGCAC	CTGATGGGAC	60
	ACTTTTTGCC	CCACTGTAAA	TTCTGGGTGT	ATCCTCCACT	GTATGCTGTC	ACCCCAAGGG	120
25	CAAGCACTGC	ATCTGCTTAG	TGAAGGATTT	ATTGTTCGGA	AGATACATIT	TCCCCTTKAG	180
23	CAGAGAGTGG	CGTATCCTGG	CAGTCTTCGG	TGAGCCAGTT	GTACCAGGAT	TATGAAATGC	240
	AGATGTTTAC	TGTGTCATTG	TIGCIGICAT	TGCTACTGAG	GAGTACTGAC	CAGAATCATC	300
30	TGCAACTYTT	AGTTGGCAGA	GAGGACCACT	ATGGCGGGTA	GCTCTTTTCT	TTCCTGCCAT	360
	TGTGGGGATG	ATTCCAGGCC	AAAGATGATG	GARAAGTATG	GAAATCATCT	GAAAGGTTGA	420
35	AGCTTGGCAC	GTGAAGCCAT	TCATGACTTT	GTAAGGCAGT	TTTGCTGAAG	GCCAGTTCTG	480
33	CCCTGGGAGG	GACGGAGGTG	AATCCTCCTG	AGTACCTGTG	GTTTTCTTAC	TTCCTGCTGA	540
	ATTTACCTAA	GTGCCTGTTG	TITGCTTGCT	GTGGAGGCTT	TCTGGTATTT	CATTTCAGGT	600
40	GCAGATGCCT	TCACTTTCCC	ACCRAAAAA	CCCCMACCAA	ACCTAAGACC	TTACTGCAAC	660
	TAAGTYTNCC	AAGTACTTTT	TAACCCAATG	GGATGAACAG	CCTGTGGTCT	GCTCAGATCA	720
45	CCCTGAGTGC	GTGTGAGAAG	GCMTNGGCTT	TGCCAGGAAA	TCCAGGAAGG	CAGGGCCGGG	780
43	CTGTGTTGGA	AGCTGGCTTA	CCTCCTCCCC	CAGCCTTATT	TCAATTAAAA	GGGCATTGAC	840
	TGGGAGCAGC	AGTCCTGGAG	TTTGTTGCAT	TTCCTATTGC	CCTCAAAATG	AGAAACCAGG	900
50	AAAATAGCAG	ATTGGAGCCT	TOGAGAAGGC	AGTAAATGGC	TGTTTTTATT	GACAAAAGGA	960
,	AAACATTTTA	CTGCCATCTC	ACTGATGGCA	TCTCACTGAC	TTAAAATGAA	GGCANGTTGT	1020
55	AGTAAAAAA	AAAGTCTACA	TTTTTCCACC	GCCACGTTCT	TATATCCTGT	TTGTCAGCCA	1080
22	CTGCTCANAA	GGGCATGTTG	TCTTGCGGAN	TANAGGCGCT	CTCCTTCCCT	CGTTTTCCCT	1140
	ATAGGTTGGG	TG					1152

(2)	INFORMATION	FOR	SEQ	ID	NO:	68:
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5 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2483 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:	
٠	AGCAGGCGGT GCGCTGGGGG CGGGAGCAGC GCGKAGCCCG GCTCGGCCAC ACCGATCGCC	60
15	CGCCGCCATG GGCTCCTCGC AAAGCGTCGA GATCCCGGGC GGGGGCACCG AGGGCTACCA	120
	CGTTCTGCGG GTACAAGAAA ATTCCCCAGG ACACAGAGCT GGTTTGGAGC CTTTCTTTGA	.180
20	TTTTATTGTT TCTATTAATG GTTCAAGATT AAATAAAGAC AATGACACTC TTAAGGATCT	240
20	GCTGAAASCA AACGTTGAAA AGCCTGTAAA GATGCTTATC TATAGCAGCA AAACATTGGA	300
	ACTGCGAGAG ACCTCAGTCA CACCAAGTAA CCTGTGGGGC GGCCAGGGCT TATTGGGAGT	360
25	GAGCATTCGT TTCTGCAGCT TTGATGGGGC AAATGAAAAT GTTTGGCATG TGCTGGAGGT	420
	GGAATCAAAT TCTCCTGCAG CACTGGCAGG TCTTAGACCA CACAGTGATT ATATAATTGG	480
30	AGCAGATACA GTCATGAATG AGTCTGAAGA TCTATTCAGC CTTATCGAAA CACATGAAGC	540
	AAAACCATTG AAACTGTATG TGTACAACAC AGACACTGAT AACTGTCGAG AAGTGATTAT	600
•	TACACCAAAT TCTGCATGGG GTGGAGAAGG CAGCCTAGGA TGTGGCATTG GATATGGTTA	660
35	TTTGCATCGA ATACCTACAC GCCCATTTGA GGAAGGAAAG AAAATTTCTC TTCCAGGACA	720
	AATGGCTGGT ACACCTATTA CACGTCTTAA AGATGGGTTT ACAGAGGTCC AGCTGTCCTC	780
40	AGTTAATCCC CCGTCTTTGT CACCACCAGG AACTACAGGA ATTGAACAGA GTCTGACTGG	840
	ACTITICIATI AGCICAACIC CACCAGCIGI CAGIAGIGII CICAGIACAG GIGIACCAAC	900
	AGTACCGTTA TTGCCACCAC AAGTAAACCA GTCCCTCACT TCTGTGCCAC CAATGAATCC	960
45	AGCTACTACA TTACCAGGTC TGATGCCTTT ACCAGCAGGA CTGCCCAACCT	1020
	CAACCTCAAC CTCCCAGCAC CACACATCAT GCCAGGGGTT GGCTTACCAG AACTTGTAAA	1080
50	CCCAGGTCTG CCACCTCTTC CTTCCATGCC TCCCCGAAAC TTACCTGGCA TTGCACCTCT	1140
	CCCCCTGCCA TCCGAGTTCC TCCCGTCATT CCCCTTGGTT CCAGAGAGCT CTTCTGCAGC	1200
	AAGCTCAGGA GAGCTGCTGT CTTCCCTCCC GCCCACCAGC AACGCACCCT CTGACCCTGC	1260
55	CACAACTACT GCAAAGGCAG ACGCTGCCTC CTCACTCACT GTGGATGTGA CGCCCCCCAC	1320
	TGCCAAGGCC CCCACCACCG TTGAGGACAG AGTCGGCGAC TCCACCCCAG TCAGCGAGAA	1380
60	GCCTGTTTCT GCGGCTGTGG ATGCCAATGC TTCTGAGTCA CCTTAACTTT GAACCATTCT	1440

	TTGGAATTGG	CGTGGTATAT	TTAACCACGG	GAGCGTGTCT	GGAAACGCAA	ACTATCATTA	1500
	ATTTCATACT	AGTTTGTACC	GTATCTGTAG	GCATCCTGTA	AATAATTCCA	AGGGGAAAAC	1560
5	TAAACGAGGA	CGTGGGTTGT	ATCCTGCCAG	GTTGAGTGGG	GCTCACACGC	TAGGGTGAGA	1620
	TGTCAGAAAG	CGCTTGTATT	TTAAACAACC	AAAAAGAATT	GTAAGGGTGG	CTTGCTGCCA	1680
0	GGCTTGCACT	GCCGTTCCTG	GGGGTGTGCA	TCTTCGGGAA	AGGTGGTGGC	GGGCCTCCA	1740
.0	CTAGGITTCC	TGTCCCCTGC	TGCTCCTTCC	GTAAGAAAAT	GAAATATTCT	ATGCCTAATA	1800
	CTCACACGCA	ACATTTCTTG	TACTITGTAA	CTCCTTTGCG	AGAATGCAGA	CCACCTCACT	1860
5	AAACTGTAAA	CGGTAAAGAG	ATTTTTACTT	TTGGTCTCCG	TGAGTCGCAT	CTCTACTAAG	1920
	GTTTACACAG	GAATTCCACC	TGAAGACTTG	TGTTAAAGTT	CTACAGCGCG	CACTGTTAAC	1980
20	TGAACGTCTT	TTTCTTCAGC	CTATACGCGG	ATCCTTGTTT	TGAGCTCTCA	GAATCACTCA	2040
-0	GACAACATTT	TGTAACTGCT	GCTGTTGCTT	TCTACATACA	CCTTATAAAG	TGACATTTCA	2100
	AAAGAAATAA	GGTGCCACAG	TTTTAAACCA	GAAGGTGGCA	CTCTGTGGCT	CCTTGTAGTA	2160
25 `	TTATAGCTAT	ACTGGGAAAG	CATAGATACA	GCAATAAAGT	ACAGTAATTT	TACTTTTTT	2220
	CTTGTGTTAC	ATCTAAATŢA	CAACCCTTAA	TTGCCACGTG	TGCACTTACT	ACTCTCCAGT	2280
30	ATGTCTTATT	ACTCTCCAGT	ATGTCACGCA	TCTTTAACTT	TTCACGTCCT	ATGTTTGCTT	2340
	TCTCCCATTT	TTAAGAGATG	GTAAGTTAAC	TGGAATTGAT	TTACTGAATG	AAATTAAATG	2400
	CAGATATCCC	TGTTTTTGAA	АТААААААА	AAAAAAAAA	AAAAAAAAA	AAAAAAAA	2460
35	AAAAAAAAA	AAAAAAAAA	AAA			,	2483

# 40 (2) INFORMATION FOR SEQ ID NO: 69:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 536 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 69:

50	GAGAAATGGA	GCTTTGTTAG	TTTAAAAATT	TTTCAACGCA	AACAGTCATT	TTCCAGTGAA	60
	AGGAGAGCGT.	ATCCGCCGTA	GGATGGACTT	AGATCGTGTA	AAAGCTGAGG	CCACCGAGGA	120
	TATAACCTCC	GGGGTCCTTT	GCCTCCTTTT	CCTTAGACTC	CCTCCAAACT	CGTGTATCTT	180
55	TCCTTCAGCA	GTACTGGGCT	CCACGCGAAC	CTAGTCCTTT	GTCTTTACCC	TATTACCTTT	240
	CATAACATCC	TAGTTGAAAA	GTARTTATTC	AACCGCGTTT	GAAAATGAGA	ACAGGTTCAC	300
60	AGARGCTAGG	TTACTTGCGA	AGGTCGTTCA	ATTAGTAACC	AGTAACGCCA	GGACTGCCAG	360

	TTTCTTGCTT CCGAATTCTC ATGGTAGCTT TCACCARGCT CCCCGTCMAA TGCTAACGTC	420
5	AACTACTGAA CTAGATTAGC AAAAAGGTCT TTTAACAGAA TTCCTGGTTT TCAGAGAGAG	480
	TTTCTTTCAT GAAGCGCCCC ATTTCTACAG AGGAAAATAA ACTCCAAGCA GCCAGT	536
10		
10	(2) INFORMATION FOR SEQ ID NO: 70:	
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 865 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70:	
20	CCACGCGTCC GGCCTTTCTT GGCCAGAGGC GCCGGTTGGA CTCACGGGCG GGGCATGATG	60
•	GGTAACAGGA CCGGTGGGGT CCCCAGGAAG TCCTAGAGGG GGTCGGGGTT TGGGTGGACA	120
25	AGCTTTCCTC GTCCTCTCCC GACAGAGCTG ACGTGTCCTG GGTTCCACCG GGAGCGGGCA	180
	TTTCCACCGG ACGGGAGGGT TCGGGGTGTC CGGGGCTGGG GAATACGTAG GGGTTGCCGC	240
30	GCGGTGTGGG GAGTTGGGCC GTGTGGCTGC AGTCCCGGGA GTTCTTGGAG GGGGTCGGCC	300
50	CACCGAGCTT CCGGACCGGC TGATCTGCCC GTAGCTTGCC GGANGGARGG CGGAGCTGAC	360
	TCTCCGTCCC TTCTCCCATC CCCTCCAGTG GTGGGTACGG GCACCTCGCT GGCGCTCTCC	420
35	TCCCTCCTGT CCCTGCTGCT CTTTGCTGGG ATGCAGATGT ACAGCCGTCA GCTGGCCTCC	480
	ACCGAGTGGC TCACCATCCA GGGCGGCCTG CTTGGTTCGG GTCTCTTCGT GTTCTCGCTC	540
40	ACTGCCTTCA ATAATCTGGA GAATCTTGTC TTTGGCAAAG GATTCCAAGC AAAGATCTTC	600
	CCTGAGATTC TCCTGTGCCT CCTGTTGGCT CTCTTTGCAT CTGGCCTCAT CCACCGAGTC	660
	TGTGTCACCA CCTGCTTCAT CTTCTCCATG GTTGGTCTGT ACTACATCAA CAAGATCTCC	720
45	TCCACCCTGT ACCAGGCAGC AGCTCCAGTC CTCACACCAG CCAAGGTCAC AGGCAAGAGC	780
	AAGAAGAGAA ACTGACCCTG AATGTTCAAT, AAAGTTGATT CTTTGTAAAA AAAAAAAAAA	840
50	ААААА ААААААААА	865

(2) INFORMATION FOR SEQ ID NO: 71:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 932 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi)	SEQUENCE	DESCRIPTION:	SEQ	ID	NO:	71:
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5	TCATCATATA	CAAAGTTTTT	CGTCACACTG	CAGGGTTGAA	ACCAGAAGTT	AGTTGCTTTG	60
J	AGAACATAAG	GTCTTGTGCA	AGAGGAGCCC	TCGCTCTTCT	GTTCCTTCTC	GGCACCACCT	120
	GGATCTTTGG	GGTTCTCCAT	GTTGTGCACG	CATCAGTGGT	TACAGCTTAC	CTCTTCACAG	.180
10	TCAGCAATGC	TTTCCAGGGG	ATGTTCATTT	TTTTATTCCT	GTGTGTTTTA	TCTAGAAAGA	240
	TTCAAGAAGA	ATATTACAGA	TTGTTCAAAA	ATGTCCCCTG	TTGTTTTGGA	TGTTTAAGGT	300
15	AAACATAGAG	AATGGTGGAT	AATTACAACT	GCACAAAAAT	AAAAATTCCA	AGCTGTGGAT	360
13	GACCAATGTA	TAAAAATGAC	TCATCAAATT	ATCCAATTAT	TAACTACTAG	ACAAAAAGTA	420
	TTTTAAATCA	GTTTTTCTGT	TTATGCTATA	GGAACTGTAG	ATAATAAGGT	AAAATTATGT	480
20	ATCATATAGA	TATACTATGT	TTTTCTATGT	GAAATAGTTC	TGTCAAAAAT	AGTATTGCAG	540
	ATATTTGGAA	AGTAATTGGT	TTCTCAGGAG	TGATATCACT	GCACCCAAGG	AAAGATTTTC	600
25	TTTCTAACAC	GAGAAGTATA	TGAATGTCCT	GAAGGAAACC	ACTGGCTTGA	TATTTCTGTG	660
23	ACTCGTGTTG	CCTTTGAAAC	TAGTCCCCTA	CCACCTCGGT	AATGAGCTCC	ATTACAGAAA	720
	GTGGAACATA	AGAGAATGAA	GGGCAGAAT	ATCAAACAGT	GAAAAGGGAA	TGATAAGATG	780
30	TATTTTGAAT	GAACIGTTTT	TTCTGTAGAC	TAGCTGAGAA	ATTGTTGACA	TAAAATAAAG	840
	AATTGAAGAA	ACACATITTA	CCATTTAAAA	AAAAAAAAA	ACTNGAGGGG	GGCCCGGTAC	90
35	CCAAATCGCC	GCATAGTGAT	CGTAAACAAT	CT			93

(2) INFORMATION FOR SEQ ID NO: 72:

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# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 996 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:

50	CGCCTGGCAC CATGAGGACG CCTGGGCCTC TGCCTGTGCT GCTGCTGCTC CTGGCGGGAG	60
	CCCCCCCCC GCGCCCACT CCCCCGACCT GCTACTCCCG CATGCGGGCC CTGAGCCAGG	120
	AGATCACCCG CGACTTCAAC CTCCTGCAGG TCTCGGAGCC CTCGGAGCCA TGTGTGAGAT	180
55	ACCTGCCCAG GCTGTACCTG GACATACACA ATTACTGTGT GCTGGACAAG CTGCGGGACT	240
	TTGTGGCCTC GCCCCCGTGT TGGAAAGTGG CCCAGGTAGA TTCCTTGAAG GACAAAGCAC	300
60	GGAAGCTGTA CACCATCATG AACTCGTTCT GCAGGAGAGA TTTGGTATTC CTGTTGGATG	360

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PCT/US98/11422

•	ACTGCAATGC	CTTGGAATAC	CCAATCCCAG	TGACTACGGT	CCTGCCAGAT	CGTCAGCGCT	420
	AAGGGAACTG	AGACCAGAGA	AAGAACCCAA	GAGAACTAAA	GTTATGTCAG	CTACCCAGAC	480
5	TTAATGGGCC	AGAGCCATGA	CCCTCACAGG	TCTTGTGTTA	GTTGTATCTG	AAACTGTTAT	540
	GTATCTCTCT	ACCTTCTGGA	AAACAGGGCT	GGTATTCCTA	CCCNGGAACC	TCCTTTGAGC	600
10	ATAGAGTTAG	CAACCATGCT	TCTCATTCCC	TTGACTCATG	TCTTGCCAGG	ATGGTTAGAT	660
10	ACACAGCATG	TTGATTTGGT	CACCTAAAAA	GAAGAAAAGG	ACTAACAAGC	TTCACTTTTA	720
	TGAACAACTA	TTTTGAGAAC	ATGCACAATA	GTATGTTTTT	ATTACTGGTT	TAATGGAGTA	780
15	ATGGTACTTT	TATTCTTTCT	TGATAGAAAC	CTGCTTACAT	TTAACCAAGC	TTCTATTATG	840
	CCTTTTTCTA	ACACAGACTT	TCTTCACTGT	CTTTCATTTA	AAAAGAAATT	AATGCTCTTA	900
20	AGATATATAT	TTTAYGTAGT	GCTGACAGGA	CCCACTCTTT	CATTGAAAGG	TGATGAAAAT	960
20	CAAATAAAGA	ATCTCTTCAC	ATGARAAAA	AAAAAA			996
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# (2) INFORMATION FOR SEQ ID NO: 73:

# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 785 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:

25	(XI) SEQUENCE DESCRIPTION. DEG 15 No. 15.	
35	GCCACGAGGG GCTTTGCGTA CACAATAGCT GCTAGGAGTA CCCAAAGCCT GARTACARCC	60
•	TECTEGTETC ATEGCCACGT GTGAGCAGGC CAGCGTCAMA CGGCTCGCTG TGACCCGTCC	120
40	CGRAGACTGA AATGGGCCTG GGTCTTCTCC TKGTCCTGTG ATWAAAGTCC TCTCTTGAAA	180
	GTGGAGAGCA AAGGCACACA GAGGTGCGCG CTCACAAGAA TTCCTCCCGG TGACTGGGTA	240
45	ATCAATGTTA CTGCTGTTTC CTTTGCAGGA AAGACCACAG CAAGATTCTT TCATTCGTCT	300
40	CCTCCTAGCC TGGGGGACCA GGCTCGAACT GACCCTGGAC ATCAAAGGAG GGATTATGTG	360
	GCTGCTAAAG CCATCGGCCC ACAGCCCTGT TCACRTCTTG GTGCTTCTCT TTCCCAGAGG	420
50	CTGGTCCCAG CCAGGCACAC ACAAAAGGCA GATTCTCGTA AACSCAGCCT CCCTCCCTGG	480
-	AGGCTGCCTC CTGCCCTGGA TCTGGAGTGG AGCTGCTCTG AGATTTTGAG TTCTTCTGCA	540
55	GAGATGATTA AATATATCCA AGAGACATTG GAAAACCTGC TGAACATTTT ACATTGGTCT	600
33	GCTCAGCACA TGGCTGGATG CGGATATTTC TATAATTCCA GAAAGTCACA CAGCTCCTCT	660
	GTATGAGACC AGTGGGCCCC ATTTAAAAGA ACAGGATGAG AATCTAAGAT ATATTATTAA	720
60	TAAATGTAAT GGATTTTTT TTTGTAAAAA AAAAAAAA	780

PCT/US98/11422

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(2) INFORMATION FOR SEQ ID NO: 74:

151	CECTENICE	CHARACTERISTICS:

(A) LENGTH: 1069 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:

15	(XI) SEQUENCE DESCRIP	IION. SEQ ID NO	. /=.	•	
	TCCTCACCAT TCCCCTAGGN CAGGTC	CCTG CAGGTCCCAC	ACTTCTCCCA	GGTCCCTAAA	60
20	CTTGGGTCGG TCCTTTCCCT GGAGTA	SCTG GNTCCTCCAG	TCGAGGTCCC	TGTTCAGTCG	120
20	GTTCTTAGGC TCCTGCACAT GAAGGT	STOT GCCTGTGGTG	TCTGGGCTGC	TCTAGGAGCA	180
	GATACAGGCT GGTATAGAGG ATGCAG	aaag gtagggcagt	ATGTTTAAGT	CCAGACTTGG	240
25	CACATGGCTA GGGATACTGC TCACTA	GCTG TGGAGGTCCT	CAGGAGTGGA	GAGAATGAGT	300
	AGGAGGGCAG AAGCTTCCAT TTTTGT	CCTT CCTAAGACCC	TGTTATTTGT	GTTATTTCCT	360
30	GCCTTTCCGA GTCCTGCAGT GGGCTG	CCCT GTACCCTGAA	CCTCATGAGC	CTCTAAGGGA	420
30	AAGGAGGAAC AATTAGGACG TGGCAA	TGAG ACCTGGCAGG	GCAGARTACA	AGCCCAGCAC	480
	CAGTGTCCCA GCCTTACTGG GTCCTT	ACCC TGGGCCAAAC	AGGGAGGGCT	GATACCTCCT	540
35	TGCTCTTCCT AGATGCCCAC CTCCTA	CAAT CTCAGCCCAC	AAGTCCTCTC	CACCCTAGGG	600
	GGCTTGCTGC ATGGCAATAA CTCATA	ATCT GATTTGGAGG	TTTGCCCTTT	ACAGGGGCAG	660
40	ATTTTCTGCT CAGTTCAACA ATGAAA	TGAA GAGGAACTCC	CTCTTTCTAC	AGCTCACTTC	720
40	TATCAGAGGC CCAGGTGCCT CAGAGC	CACA TIGAGTIGCI	TTTTCTGGGA	TGAGGAAGTA	780
•	GGGTTAAACT CCCCAGTTTC CTGAGG	GAGG CTCCTGACAG	GTGCCCTTTG	TCAGACCCTA	840
45	CCACAGCCTG GATAGGCAGC CACATT	GGTC CTCGCCCTTC	CTCGGNACTC	CGTGGTGGTC	900
	CTGCCCTTCT CCCTGCATGC CTGTGC	GTCT GCTCTGGTG1	GTGAAGGTCG	GTGGGTTAAC	960
50	TGTGTGCCTA CTGAACCTGG CAAATA	AACA TCACCCTGC	AAGCCAAAAA	AAAAAAAA	1020
50	AAAAA AAAAAAAAA AAAAAAAA	AAAAAAAA AAAA	AAAAAAAA	•	1069

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(2) INFORMATION FOR SEQ ID NO: 75:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 831 base pairs

(B) TYPE: nucleic acid

WO 98/54963 PCT/US98/11422

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	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 75:	
J	GGACATTAGA TCACTGTGGA CCTAAAACAA ACAAACAACT ATAAGGAAAA TGGCATTAGA	6
	AATGGTCTGG GGATCAGTTT ATCACTGCAG TTGTTACATC ACCCCATGGT CTAAAATACA	120
10	GAGCTTTAGT CTGTCTCTGT TTCAGTTCAT TTTACAGGAG GTGAACATCA CACTTCCAGA	186
	AAACTCTGTC TGGTATGAAA GGTATAAATT TGATATTCCT GTCTTTCACT ŢGAATGGCCA	24
15	GTTTCTGATG ATGCATCGAG TAAACACCTC AAAACTTGAA AAACAGCTCC TGAAACTTGA	30
13	GCAGCAAAGT ACTGGARGCT GACTGATGCC CTCATGATTT TCCACCCTCT CTTCCCATAA	36
	AGCATCTTCC TAAGGAAATG AMCATGGCCT GATACTCATT TTGTCACTTG TACAGAGCCC	42
20	TAAGGATGTT CTGAATTCAG TGGTGCCAAA TAAATGTTGA CATTCCCCTT TTGGTTGATG	48
	GAAGTATCAG TGTGGGAACT GTTTGCTTAA TGGCATTTTA TAAAATAAKA AKAKCATATT	54
25	AGCAGGGAGG GAGATGATGG AGGGAGGGAG AAGTCCATTT GTCTTATTTA TCCTTTTTGT	60
23	ATTAATAGAG AAGCACTTCA CAGTCACTGG CAATGCCATT TATAGGAAGA AGGTTCTGCA	66
	TICCTGCTGC TCCCGGAGGG CTTAACTTTT TAATGAAAGA ATAAATGCTC TTCCACTCAG	72
30	TAGATAAAGT GAAATGTGAA TTGTTAATAA CTGTGCACGG TCAATAAAGC GATGTTTTAA	78
	GGAATACAAA AAAAAAAAAA AAAAAAAAAA AAAAAAAAA	83
35		
	(2) INFORMATION FOR SEQ ID NO: 76:	
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 590 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:	
	TATATATAGA CNGTTAATAG TOGTGANTGN TGTGNACGAA CATTAACGGA AGTAGCATGT	6
50	AGCCAGTCGA ATAACNTATA AGGACAAAGT GGAGTCCACG CGTGCGGCCG TCTAGACTAG	12
50	TGGATCCCCC GGCTGCAGGA TTCGGCACGA GCTGCCAGGT GAGGAGCAGA GAGACTGTTC	18
	CCTTGGGTGG AGAGGTGTGG GCATGAGAGC CACCCATTGC CAAGCAGCAA GAATGTTCGT	24
55	COMPANIENCE CHARCES SAN SARCESCOCHE CACCOMICCA STACCOCCOCC ACACAGCATATA	30

GCTTGTGTTT CTCCTGTCCC TGTTCTCCCG GAGGGCCCAG GTGGAACTCA CGACAGGGAG

GGAGACGCTT CCCAAAAACC TGCAGGGCTA TTTCCCAGAA TTTGGTTTTC AAGTACAAAA

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WO 98/54963 PCT/US98/11422

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	CTTTTTGTCC	TGTAAGATAT	ATGCAGCCTC	ACAGAAGCAG	CCTCTGCCTC	CACTTTACCA	480
	GCTACGTTTT	TATCTTAAGC	ACATGGGGCT	CCCTTAGAAC	TTACTCCACT	GATTTAAAAA `	540
5	АААААААА	AAACTCGAGG	GGGGCCCGG	TACCCATTCG	CCCTAAAAGT		590

10 (2) INFORMATION FOR SEQ ID NO: 77:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1274 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:

		-		-			
20	GAGCCACCAC	ACCTGGCCTG	GAAGGAACCT	CTTAAAATCA	GTTTACGTCT	TGTATTTTGT	60
	TCTGTGATGG	AGGACACTGG	AGAGAGTTGC	TATTCCAGTC	AATCATGTCG	AGTCACTGGA	120
25	CTCTGAAAAT	CCTATTGGTT	CCTTTATTTT	ATTTGAGTTT	AGAGTTCCCT	TCTGGGTTTG	180
<b>2.</b> 3	TATTATGTCT	GGCAAATGAC	CTGGGTTATC	ACTITICCTC	CAGGGTTAGA	TCATAGATCT	240
	TGGAAACTCC	TTAGAGAGCA	TTTTGCTCCT	ACCAAGGATC	AGATACTGGA	GCCCCACATA	300
30	ATAGATITCA	TTTCACTCTA	GCCTACATAG	AGCTTTCTGT	TGCTGTCTCT	TGCCATGCAC	360
	TTGTGCGGTG	ATTACACACT	TGACAGTACC	AGGAGACAAA	TGACTTACAG	ATCCCCCGAC	420
35	ATGCCTCTTC	CCCTTGGCAA	GCTCAGTTGC	CCTGATAGTA	GCATGTTTCT	GTTTCTGATG	480
<b>J</b> J	TACCTTTTTT	CTCTTCTTCT	TTGCATCAGC	CAATTCCCAG	AATTTCCCCA	GGCAATTTGT	540
	AGAGGACCTT	TTTGGGGTCC	TATATGAGCC	ATGTCCTCAA	AGCTTTTAAA	CCTCCTTGCT	600
40	CTCCTACAAT	ATTCAGTACA	TGACCACTGT	CATCCTAGAA	GGCTTCTGAA	AAGAGGGGCA	660
	AGAGCCACTC	TGCGCCACAA	AGGTTGGGGT	CCATCTTCTC	TCCGAGGTTG	TGAAAGTTTT	720
45	CAAATTGTAC	TAATAGGSTG	GGCCCTGAC	TTGGCTGTGG	GCTTTGGGAG	GGGTAAGCTG	780
45	CTTTCTAGAT	CTCTCCCAGT	GAGGCATGGA	GGTGTTTCTG	AATTITGTCT	ACCTCACAGG	840
	GATGTTGTGA	GGCTTGAAAA	GGTCAAAAAA	TGATGGCCCC	TTGAGCTCTT	TGTAAGAAAG	900
50	GTAGATGAAA	TATCGGATGT	AATCTGAAAA	AAAGATAAAA	TGTGACTTCC	CCTGCTCTGT	. 960
	GCAGCAGTCG	GGCTGGATGC	TCTGTGGCCT	TTCTTGGGTC	CTCATGCCAC	CCCACAGCTC	1020
55	CCAGGAACCT	TGAAGCCAAT	CTGGGGGACT	TTCAGATGTT	TGACAAAGAG	GTACCAGGCA	1080
33	AACTTCCTGC	TACACATGCC	CTGAATGAAT	TGCTAAATTT	CAAAGGAAAT	GGACCCTGCT	1140
	TTTAAGGATG	TACAAAAGTA	TGTCTGCATC	GATGTCTGTA	CTGTAAATTT	СТААТТТАТС	1200
60	ACTGTACAAA	GAAAACCCCT	TGCTATTTAA	TTTTGTATTA	AAGGAAAATA	AAGTTTTGTT	1260

TGTTAAAAAA AAAA 1274

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### (2) INFORMATION FOR SEQ ID NO: 78:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1133 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:

	AGGATTTTTC	CTTGTTCAAC	CAAAATCTGA	GCATTCTTTC	TATGTTGAAA	ACACTGAAAA	60
	ACTAATTTWA	GTTAATGAAC	TAGAAAGAAT	ATTGATTTTW	AAGAAACAGA	AAAATACTAC	120
	TTATTTTCCT	TCTCAAATAA	CGTTTCTTTC	AAAAACTTCT	GGCTGAAGTA	TAACATGCTG	180
	GTAGTTAACA	TAAATCTTGT	CTTTCTCTTG	TTCTTTATCT	TTCTTTGTTA	TTTAGATGCT	240
	TGTATAAATG	TCTTTTGTTT	TTATTAAGTG	CCTAATTGAC	AGAGCTTAAT	TTGAAGAAGT	300
	GCCCTAATTT	ATTGACCACT	TAAGAATTGC	CTTTATTGGG	GTATTTTATT	TGTTCCTGCG	360
	TCTTTTTGAT	GTTGTTCAGT	CTACTCATCC	CTGTGAGTAT	GTGTGGGGGA	CAGCTGATAG	420
	AAGGGAGGAG	AGTGTGTCTA	TGCTCAGGAT	TGCCCTTTAG	CCACTCAGCC	AGAGATCCAC	480
	AGGGAGCAAC	AAGGACAGTT	TCACATGCTT	AGACTTTCTT	GGAAGAAACA	GTGAGGAGGA	540
	GTAAGTCGTG	AGTAGTGTCA	AGCTGGATGT	AGAATTGTCC	TAAGGCAGTT	GACCCCACCT	600
	TCCAACATGT	TTTCACTTTA	TTTCCCCCTC	CCTACATTTG	GGTTAGGTTC	CATTTGGATT	660
	TGCAGCAATA	ATGACTTTAT	TTCTCTCTTG	GTCAGGATTT	GGCACATAAA	ATCCTTTTAT	720
•	TATAGAACTA	GCTATTTTAG	TTACATAGTA	ATGTAACTAA	TGGAGAGATT	TATAGAGAAT	780
	TTTGKTTTTG	CTGTCATATA	TGTCCATTTT	GGAGACAGAT	ATGATAGAAC	TAGAAATTAA	840
	GTTGCATTTC	TGCAAGTGCC	ATTTGAATGA	ACTTCAAGTA	TCTTCTTAAT	TATTAAATTT	900
	TCTGATGAAG	GCATTGTAAC	AAATATATAG	TATTATTAAA	TCTAATTAAT	ATTTGGAAAT	. 960
	ATTAATAAAT	AGGTATTTTA	TTTACTGTAA	AAAGTCAAAC	TTCATTATGT	AGATAAATCT	1020
1	TATTCTTTTC	ATTCTTTCCC	CTGTTTACAT	CCTTTTTACA	AAGCTTAGTC	ACCAATTAAA	1080
	GCTTTCCTAT	СААААААА	ААААААААА	ACTCGAGACT	AGTTCTCTCT	CCT	. 1133

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### (2) INFORMATION FOR SEQ ID NO: 79:

60 (i) SEQUENCE CHARACTERISTICS:

	<ul><li>(A) LENGTH: 661 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li><li>(D) TOPOLOGY: linear</li></ul>	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:	
	GAATTCGGCA CGAGGGGAAA AGGATGCTGA ACGAGAGCAG AAAGCCTCTT TCCTTTGCTT	60
10	CACGCCTTTC CAGTCTTTAT TTTAAACTCG GGTTCCCTTT CTGTGGTCGC AGCAACCTTT	120
	ACTCCACCTG CACTGCTGCT CCTGGGGGCT CCCCAGGCCT CCCTCTGCCT TTCTACCCAG	180
15	TGGCTGACGG GATGCCTGTC TTGCCTGGAC GCACCACTGC TCTCCTGTCC CTCACCTTGG	240
15	CTTTTGCTGT GCCCTGCTCT GGGGTTGAAG CTGGCCCATG TGTCCCCCGG AGTCATGGCT	300
	GCTCCTCCTG GGAGGCCTCT GTGTGCGTCA CGTCTTCCAC ACCTGGGGGC AGCTGGCGAG	360
20	CCCGTGCTCT GTTCCCCTCG GCTGCTTGGC ACAGAGYTGC AGCCTGGGAY TCTCCGTGGA	420
	CCCAGACTGG GGATTTTGCC AGGGGGGCGA TGGGAGGAGC AGGTGCTTTG CCTGGCGGCT	480
25	GTGTCTGCAT TTCTGGACGC CCCAGAGCAC AGAAGTTGCC GGCACTTTGA GGTCTTCCTC	540
	GGCATGTGCC AGATTACATG AGTGACGGCT GGGAATATGT TTTCTTTTTT GTAATGGAGG	600
	CGTGTTTCAC ATATAGTAAA GCTCACCAAA AAGTAAAAAA AAAAAAAAA AAAAAAACTCG	660
30	A	661
35	(2) INFORMATION FOR SEQ ID NO: 80:	
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1378 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:	
45	ATTGGGTACC GGGCCCCCC TCGAAGTTTT TTTTTTTTT TTTTAATGAA AGCTCTCAAA	60
	TAAGCGATTT TATTCCTATC CATGATTGCA GACATTTACA AAACCATAAC ATCTGAGTTC	120
50	ACCTTAAAAA ATAACTTATA TAAAGCAGTG ATATACACAG CACAAAATAG TTCAGGGAGG	180
50	GGGCAGGAGC AACTTGTAAT AATTAAAATG TAAACGTGAA AAAAAGGATG GAATAAAAGT	240
	CCCTACTTAT TICTACTTAA GATGTCATGT GATAATATTT TACAATGTCC TGTGGGTCAA	300
55 .	TGTATGTATG TGTATATGTC TGTATAACAT ACACATATAC AGTACATTCT CTTTCCCACA	360

CGTACAGTTG TTTGAATCAC ATTTGGACCC GCTTTCTTCA CAAAAGAGGG GAGAGAGCAG

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	GAAATAAAAA	GGTTGGTTTG	GTGTGACTGA	GATTCCTTTG	TTTAACTGTA	CACTGTGATG	540
	AATAATTTTC	TTCCGTAGTA	GTTCTGTGAA	GGGCTGACTC	ACTGTGGTTT	TCATGAGGAG	600
5	ACTTGGTAAT	GGATCACACG	CTCATTGTCA	TGCTAGGGGA	GTAACTCTCA	CTCTGAAAAG	660
	GATTTAAGAA	ATTTCCCCCC	ATTTCGCCAT	CATCCCTTGG	AGTGCCCGGT	TGATTACTCA	720
10	GGCTCATATT	ATTGGGAGAA	TTCTTGGAAA	TACTGTCCAT	ATCTCCTGAG	CCTAAAGAGC	. 780
10	CATTCATGTG	ATGTGACTCC	ATTCCTCCTA	ATCCACCCAT	GGGACCATCT	GACCCAGGRC	840
	CCATTGGAAA	ATTAGGTCTG	TTAGGTCCAG	GAGGTACTGC	ATTCATTAAA	GTATACATGT	900
15	TATCACCAGA	GTTGGTTGAA	TCTGCTGGAC	TAGGCATGAT	GGGTGTTCCT	GGTGGCCCTC	960
	CACCTCCTGG	AGGACCTACA	TAATTCCCAG	GAGATGCTGA	GGAGTATGGT	ATTGAATTGG	1020
20	CATTTGTTGG	GTTTGGCCAA	GGTCTACCAC	CACCTGGACC	CATGTTCATT	CCAGGCATTC	1080
	CAGGGCCACC	TAAAGCATTC	AGTGGGGGTC	TCATTGCACC	TCCATAGTTC	TGTGGTCCTA	1140
	AGGGCACCAT	TCCTCTTGGA	GGAGTCATTC	TCTGCATTGG	CCCACCCATA	TTTGGATGTC	1200
25	CTTGTTGTCG	AGTTGGATCC	ATTCCACTGG	GGAGTAATGG	CTGACTTCCT	GGGACACCTC	1260
	CAAGTGCCTG	ATTAGGTATC	CTCAATGGGG	GCCTTGGACC	TCCAGGGTAC	CGAGGTGACA	1320
20	TAAAAGGGTA	ATCATGGAAG	GCTTTTGCTT	CACTTGAGTG	TTCACATGTT	TCACGTCT	1378

### (2) INFORMATION FOR SEQ ID NO: 81:

35
(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1440 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

40 (D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:

ACTITIGICCA AATGIGICIG TCACATGIAG TCAGCIGNAG NAATITAAAA TGAATIGCCA 60 45 AGTGAAGAGT CTGTGGATTA ATTGGCCGTT AATTAACAGG CTTTATCAAT GTGTCCTCAA 120 180 GGGAGAGGCC CAACCCTAAT TAAGGAGCTA AACTTCCTGA GTGAGGGGCT GTGAGGATGG 50. AGGTGGAGGA GGCATCTGGG GCGGGTGGTG GCCGGGCCAG CAGATGGCGC CTCCCTGGCT GAGCTGCCCG CACCGCCAGT TCCCTCATTT CCACTCAGGA AGGCAGAGAA GGCAGAGTGA TCTCCTCAAG GAAGAGCTTC CCCAGCCTTC GGGAGCAGCT GGCAGGGCGT CCGGGAATAA 360 -55 GCCCTACACG CCGCCGCCTG CCTCCAACTC ACTAACCCTG CGCCTCTTGT CTTTCAGATT 420 CAACGCGTTC AACAGAAGCC ATCCCCAGCC CAGCTTAAAT TATAAAGATA GACAATAACT 480 CTGTTCCAAT CTGCGTGGTG CTTCTTTAGT AAATACTGTA CAGATTTTAC CATGGAGAAC 60 540

PCT/US98/11422

	TTTTTTTTTA	GTTTTTACCT	TTTCTTAATT	ACCCTTATTC	CGAATGGACG	AACACTTTCT	600
5	ACCACTGCTG	ACCATTGTAA	AATACCGTGT	ATATAAATCC	CATTGAAATA	ATGCCCTGGA	660
J	ATAGAACATC	TCAAATGCTG	CTTAATTACA	GACTCAGGTC	GATTACTTGT	ATTTCATGTA	720
	ATGTTCCTCC	AAGTTAGACA	TCTGGTGCAA	GACCAACCGG	GAGACCATGG	AATTGTCAAA	780
0	AGTACAAACT	GACAGTGTGT	ATATTTAATT	TAAAGACTTA	TTTAAAAACT	CACAAGCTCT	840
	CACCTAGACT	TTGGAGAGCA	GTCTGTTTTC	TGTAATGTCT	GATACTAGAA	ACTAATTTGC	900
15	TTATTTTAGT	TGTATTCAAG	ATTTGAAGAT	GTATTTTATA	GACAAGTTCT	GTTTTTGAAC	960
IJ	TTTGTGGAAC	TGTTCCAATC	AATCAATTTC	CCAGTTATGA	TGAGTATTTA	CATTATGAAT	1020
•	GTATAACCCA	GACATGATTT	GTAAAGCCGA	CAGTATGTTT	CTATTACACA	ACACTTTTTG	1080
20	ATACAGCGTC	TCTTCTCTTC	ACTGATACTG	GAGTCTCCGT	TGTCTGCNNG	GTCCCTTCGA	1140
	GTTTCTAGTT	ACAGACACAA	TCATACTGTG	ATTTTATTTT	TAATATGGAT	ATGCTATCAA	1200
25	ACTGTGATAC	ACTTATAATT	CACTGGTCCT	GCATCAGGAG	ATGGAGTGGG	GAAÁACTGTA	1260
43	TTTAATACAG	TTTGTATCTG	AATAATCTGT	ATGGTTTATA	CAGTTTGTGT	TGTTCAGAGA	1320
	TGTTTAAAGT	TTGATCTTTG	TTTTTCTAAA	GATTAAAAAA	GCACTTGCCC	CACTGTAAAT	1380
30	ATACAGCATG	TAAAATTICT	RTAGTATATA	AATGGCAGCA	AATCACAAAA	AAAAAAAAN	1440

# 35 (2) INFORMATION FOR SEQ ID NO: 82:

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# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1381 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 82:

45	CCCGGGCTGC	AGGAATTCGK	YACGAGGCCA	GCAGTTGCTC	CCAGTTCAGG	AGGTGCTCCT	60
	GTACCCTGGC	CACAGCCCAA	TCCTGCCACT	GCTGACATCT	GGGGAGACTT	TACCAAATCT	120
50	ACAGGATCAA	CTTCCAGCCA	GACCCAGCCA	GGCACAGGCT	GGGTCCAGTT	CTGACCTGAG	180
30	CACGGTTTTT	CCTCATGTGA	CTTCTGGGAA	GCCCTCCCT	CATCTGGGCC	AAAGGAAGGA	240
	GGACGAAGCC	CTCCTCAGCT	GCCTGTGTT	TGGGGCATGA	ATCTCTCCTC	TCCTCCTTGT	300
55	CTGGCTCTGT	TGACAAACCG	GGCATGTTTG	GCAGTAAATT	GCACCGTGT	CACACTGTTT	360
	CCTGGGATTC	AAGTATGCAA	CCAGAACACA	GGAGAAGAAA	AGCTCCAGGA	TCCCTGTCCC	420
60	CATCTGTCCT	CTTGATGTGA	GAGAGACTCT	GAGACTTCTT	CCATCGCAAT	GACCTGTATT	480
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	AAACACAAGC	CCCCCAAGCA	AAAGAAGAGG	TTGAGTTTGC	TGCCAGGATT	CAGATCAGCC	540
	CTTCCCAGGG	TCTGCAGGTG	TCACATGATC	ACAGTTCAGC	GGGAGGCTTT	CCGTACCCAC	600
5	ACTGGCTGTA	GCACTTCAGT	CCATCTGCCC	TCCAGAGGAG	GGTTTCTTCC	TGATTTTTAG	660
	CAGGTTTAGA	GGCTGCAGCT	TGAGCTACAA	TCAGGAGGGA	AATTGGAAGG	ATTAGCAGCT	720
0	TTTAAAAATG	TTTAAATATT	TTGCTTTGCT	AATGTGCTGA	TCCGCACTAA	CTCATCTTTG ·	780
U	CAAAAGGAAC	TGCTCCCTCG	GCGTGCCCCA	GCTGGGGCCT	CTGAAGGGAT	TCCTCACTGT	840
	GGGCAGCTGC	CCTGAGCTTC	AGGCAGCAGT	GTTCATCTCT	GGCCAGTTGT	CTGGTTTCCA	900
5	TGTATTCTAG	GCCAGGTAGG	CAACACAGAG	CCAAGGCGGG	TGCTGGAAGC	CAGACGGAAC	960
	AGTGTTGGGG	CAGGAAGGTG	GATGCTGTTG	TCATGGAGCT	GTGGGAGTTG	GCACTCTGTC	1020
20	TGCTGGTGGC	CCTCTCGGCT	CACATGTTCA	CAGTGCAGCT	CCTGGCAGAC	TTGGGTTTTC	1080
20	TCTTTGGTGG	TTTCTAAAGT	GCCTTATCTG	CAAACAACTT	СТТТТСТССТ	TCAGGAACTG	1140
	TGAATGGCTA	GAAGAAGGAG	CTCAGTAAAC	TAGAAGTCCA	GGGTTGCTTG	GTTTACTGGT	1200
25	TTATAAGAAA	TCTGAAAGCA	CCTCTGACAT	TCCTTTTATT	AACTCACCTC	TCAGTTGAAA	1260
	GATTTCTTCT	TTGAAAGGTC	AAGACCGTGA	ACTGAAAAAA	GTGTTGGCCT	TTTTGCGGGA	1320
30	CCAGATTTTT	AAGATAAAAT	AAATATTTTT	ACTTCTGTCA	АААААААА	TATAAAAAA	1380
<b>)</b>	C,		•				138

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### (2) INFORMATION FOR SEQ ID NO: 83:

#### - (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1706 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83:

45 ACTGCACCAC TGCCCAGGTC TCCCGGCTGG ATGAAGACGT GGTCCATGAG GAAGCTGGCT 60 AGCTCAGACT GGAGAGTAGC TTCAGGAAAA AAGACAAGTG GCCTAAGGAA ATCACGGCCC 120 50 CCAACTATCA TCTGAGGGCT AAAGATGAGA AGTAGATCAC TTAATAAGAC AAAAGCCTGT 180 AGGGGGAAAA GAAAGGATGT TTAAAAGGAC AGAATGTTTC CCAAGGTAGA AATGACACTG 240 TCAATTTCTC CTTGGAATGG GGGCAGGGAT ACTCGCCTTG TTGCTCCCAC TTGAGTCAGT 300 55 ACTCACCTGC TCCTGGATCT CAGTATCCAC ATCTGAGAGG CAACTCTGGC AGAGTTCACA 360 GAAGGCCACC ATTCTGTCCC TCAAACTCGA CAGCTGCTTC TGTGGGCACA GTGGCTTGAA 420 60 GGGGAAGAAT GAAGACACAG ACTCCTCTGT TCCCATTATC CCATCTAAGA CCCACACTCA 480

	CCTGGGGAAG	CATCTGATTT	AGAAATGTGG	GTTAGTGTCC	AGAGAATGGA	AAAATAGACA	540
5	AGAGTCAAGG	CTGGCAGGAT	AACCTGTAAC	AACAAAGGGT	TTGAAAAATG	AGGTTTGGGT	600
J	TAÇGAGAGGG	AGAGACAGAT	AGCCAGAAAC	ACACCAGTGA	AGAGGAGAGA	AAATGAGTAA	660
	AGGGAGAGCT	AATTCCTTTT	CCAGTGGAAA	ATGAGTGATA	TTCTGGACAT	TCTTCAGAGG	720
10	CATCTACACG	AAGTAGAAAT	GTCACCGCTC	CCTAATTTAC	TCTACGTCTT	CTAGAATCCC	780
	TCAATATTAT	CCTTGGCTTC	CAGGAAATCC	AAGAAGACCC	TGGAAGTAGA	GTCCACCTTC	840
15	TAAGAGAGGA	ATGTAAGAGG	TGACCCCCAC	CCACCTGATC	TTCCTCGCTT	TGTCCACTCC	900
IJ	ACGCACTGAG	ACTTGACACA	CCTAGTGGCC	ACCTAGAACG	TAGGTCCTTA	AAATYTAGCC	960
	CCCCAGCCCC	CAACCCATCT	CTAGCCTGTC	CACTCACCTG	GTGAGGAACY	TYTCCTGTGT	1020
20	CCACAGCYTT	CTGCAGGAGT	TGGCAACATG	GCTCATAGAG	CTCCCAGCGA	GTCAGGTCAT	1080
	GAGTGCTTTG	GGGGAGAAAG	GGGAATGTTA	TACTGGAAAA	GAACAGAGGG	AACCAACTCC	1140
25	ACAGACACCA	GTAAAAACGG	GATGGGGAAG	AGGAGGAAAG	CCACTCACTT	GTAGAAGGCA	1200
4.5	GAGAGGCGTT	TCAGAGTGGC	TGCCAGATTA	TATACCTCAT	CCTCATCTAG	GAAGGACGAC	1260
	TGAGAAGGAA	AGAAGATCCA	CAATAGCATT	TCCCCCAGAA	CTCATCAGTC	CACATCCCCC	1320
30	GTCTTGCAGC	CCCTCCCACC	CTTGTTTGGG	GTGTCCCATT	GTCCAGCCCC	AGCTCCTACC	1380
	TGTAACAGCT	CTTCAAGCTC	CTGCTGGAAR	CGGTCAGTCA	GCAAATCTAC	TAGCTGGCTG	1440
35	CGGGCAAAGT	CCGCCCGGCT	GAAGAAAGTG	AATTCGGGAT	TACAGAGCAG	GTAAGAGCAT	1500
	GCGCCCCAGC	CTCAAGCACC	GCTGGCTCTG	CATGCTTCAC	CACCACCTCC	TGGAGTTGCT	1560
	GCAGGAACAG	CTCCAGGTGC	TGAGAAGAAA	AGGCAGAAGA	TGGTGTGCTG	TGGGGATGGG	1620
40	AGGAGGACAC	TCTTCTGGCG	GGAAGTGGAA	CGGGGTTAAA	AGCATTAAAC	TTCAAGGATA	1680
	AGATGCCTAA	. калалалала	AAAAAA .				1706

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### (2) INFORMATION FOR SEQ ID NO: 84:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 573 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:

GAATTCGGCA CGAGCTTGGT AGCCTTAGAA CTGCATGAGC TGCTTTACCA CTGGGAAACA 60
CGAGCACAGC CTAGCTTGAT TTTGTATGTG GTATCAGATC TAAGGTGGAT GGAATTCAGG 120

	ACTTCCTGTC	TACTCTTTGA	TTTTGTTTTA	TTTTTAGAAA	TGTTTTATTT	TGTTTTATTC	180
	ATTTATTCAT	CTTCAGAGAC	ATGGTCTGGC	TCTGTTGCCC	AGGATGGAGT	GCATGGTGTG	240
5	ATCATAGGCC	ACTGCAGTGT	TGAGCTCCCG	GGCTCAGGCG	ATCCTCCTGC	CTCAGCTYCC	300
	TTAGTAGCTG	GGACTATAGG	CACATGCCCT	ACCATGCCTG	GCTTTGTCTA	CTTTTTGAAT	360
10	GATGTCYCAA	ACTAGAAGGT	CTATTAATTT	AAAAAATTAA	GGATAGCATG	CCATAATTAA -	420
10	AAATAATAAC	AGTGGGAAAA	GGCACCTTCC	AATGATTCAG	ACATCAACTT	GTGATTTAAA	480
	AAAACGAAAA	ATAAATAATA	GGAAAAAAAG	GGGAAAAAGT	ТАААТАААА	TAAAATTAAA	540
15	AAAAAAAA	AAAAACTCGA	GGGGGGCCCG	GTA			573

# 20 (2) INFORMATION FOR SEQ ID NO: 85:

### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 684 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 85:

30	CTCTTTGGCT	GTGTCTACCT	CCTTCATCTG	CTGCGCCGAC	ATAAGCACCG	CCCTGCCCCT	60
	AGGCTCCAGC	CGTCCCGCAC	CAGCCCCCAG	GCACCGAGAG	CACGAGCATG	GGCACCAAGC	120
35	CAGGCCTCCC	AGGCTGCTCT	YCACGTCCCT	TATGCCACTA	TCAACACCAG	CTGCYGCCCA	.180
33	GCTACTTTGG	ACACAGCTCA	CCCCCATGGG	GGCCGTCCT	GGTGGGCGTC	ACTCCCCACC	240
	CACGCTGCAC	ACCGGCCCCA	GCCCTGCC	GCCTGGGCCT	CCACACCCAT	CCCTGCACGT	300
40	GGCAGCTTTG	TCTCTGTTGA	GAATGGACTC	TACGCTCAGG	CAGGGGAGAR	GCCTCCTCAC	360
	ACTGGTCCCG	GCCTCACTCT	TTTCCCTGAC	CCTCGGGGGC	CCAGGGCCAT	GGAAGGACCC	420
45	TTAGGAGTTC	GATGAGAGAG	ACCATGAGGC	CACTGGGCTT	TCCCCTCCC	AGGCCTCCTG	480
43	GGTGTCATCC	CCTTACTTTA	ATTCTTGGGC	CTCCAATAAG	TGTCCCATAG	GTGTCTGGCC	540
	AGGCCCACCT	GCTGCGGATG	TGGTCTGTGT	GCGTGTGTGG	GCACAGGTGT	GAGTGTGTGA	600
50	GTGACAGTTA	CCCCATTTCA	GTCATTTCCT	GCTGCAACTA	AGTCAGCAAC	ACAGTTTCTC	660
	TGAAAAAAA	ААААААА	AAAC	•			684

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(2) INFORMATION FOR SEQ ID NO: 86:

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 1036 base pairs

(B)	TYPE: nuc	leic	acid
(C)	STRANDEDN	ESS:	double
(D)	TOPOLOGY:	line	ear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 86: TGGAGGCAGA TGCACAGGAG AAAGGTTCCC GTCCGCACCC TCTCAGACCT GAGGCTGAGC 60 TTGCAGTGAG GGCTTCTCCT CGGCCCCTCG CCCGCCCCCA GAGCTGCCAT CCCTGCTGTT 120 10 ACAAGCCAGA GGAGCCCGGA TGTGAGGCCC CAGATCACCT CCAGGGACTT GGGGTTCCCA 180 TCTGAAATCC TTTATTTTG TACCATGGGG TGGGCCCCGG GCTGAGAAGG AAGAAGCACC 240 15 CTCTCCCCGG CCTCCTCTGT CTGCACCCGT GGGGCTGTGA CTTACTCCTG CCTCCAGGGG 300 CGGGGCGGGG CCCCCTGGGA CCTCTTAAGG CCCAAGGTGG GCCCCAGGAC CTYTGGGCAG AGTGGAYTGC TCATGGCAGA TGTGTGGCAA TGTCTGGCTG WGTCTTTCCG GCAMCTGCGT 420 20 YCCCTYTCCC GGGYTCCCCT GCTGCATGGT GGATGTGCTC CTTCCTGGCC CGGTCACATT 480 GCCTCCTTGA GCCTTAGTCC AGGGGGTCAC TYCTCCCACC CCACCTACCT CACAGGGTTG 540 TTGTGAGGGT GCACAGAGGA GCAAAGTCCC TGAAGGCCCT CAGGCAGTAT ATAGGGGCCG 25 600 CCCACCTTCA GCTGCCCTGG GATGGGAAGG ACCCAGCCCG ACCCCTGGGC ATAACACTGT 660 CTTTCCAAAT GGAGATTCAG GTATTGGGGA TGCAGGTTGT GGGGAGCTGG CCTGGCAGAG 720 30 TAGGGGTAGT TGGCTTGGCC TTCTCTTTGG TGATCCCACC CCCAGCCATT TGCATTGCTG 780 GCCCAGCGC TGGCCTGGGG GGCGGGAGA GGCAGCAGAA GGGGCTGGGC AGGGGCGGTG 840 35 GAGGACTCAG GAACTGCCCG GGGAGAGTGG GTATGGCGGC TGAGCCAGGG GCCCTCCTGT 900 GTTTGACTTC CCGGGATGGG TCCTTGCTTC TCAGCTGTGT CCGACCCCAC CATGTAATAA 960 1020 40 1036 CCCNGGGGGG GNCCCG

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#### (2) INFORMATION FOR SEQ ID NO: 87:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 908 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 87:

TTAAACAAAT GGAATCATGC AATATGTGAC CTTTTGCGTC TGGCTTATTT TATTTAGCAT 60

AATGTTTTTG AGGTTCATCC AAGCTGTAGC ATGTATCAGC ACCTCATTTC TTTTTCTGGC 120

TGAATATTAT TCCATTATAT GGATTTACCA CAATTCATTT ACCTATTCAT CTTTTGTTTC 180

	TGCTGTCTGG	CTATTGTGAA	TAATGCTTCG	ATAAACATTC	ATATACAAGT	TTCTATGTGG	240
5	CTTTATGTTT	TCATTTCTCT	TGGCTATCTA	CATGGGAGTA	GAATTCTAGG	TCATAATATA	300
3	ATTTTATGTT	TAACTTCTCA	AAGAATTGCC	AAAAGGTTTT	TCATAGTGGC	TGCATCATTT	360
	ACATTCCCAC	CGGCAATGTA	CAAGGATTTC	TATTTTTCCA	TATCCTTGCA	CTTACCAACA	420
10	CTTCTTTTTK	GTWATWATTT	TGTTTTTCA	TTATTGCCAC	CCTAGTGGAT	GTGAAATGGC	480
	ATCTTATTGT	TTTGATTTGC	ATTTCTCTAA	TGACAAATGA	TATCATACTT	TTTTTATGTG	540
15	CTTACGGATC	AAAGGTATTT	CCTTGGAGAA	ATGTCCCTTC	AAGTCCTTTG	CCATTTCAAA	600
13	ATTTGGTTAT	TTGTCTTTTA	TTATTCAGTT	TTAAGAAATT	CTGGCCAGGC	GCAGTGGCTC	660
	ACCTGTAATC	MTAGCACTTT	GGGAGGCCAA	GGCGGGCAGA	TCACTTGAGK	TCAGGACTTC	720
20	GAGACCAGCC	TGGCCAACAT	GGTGAAACCC	CATCTTACTA	AAAATACAAA	AATTAGCTGG	780
	GCGTGGTGGC	AGGTGCATGT	AATCNTATCT	ACTCÁGGAGG	CTGAGGCAGG	AGAATCGCTT	840
25	GAACCCAGGA	GGCGGAGGCT	GCAGTGAGCC	AAGATCACGC	CATTGCACTC	TAGCCTGGGT	900
25	GACACAGA						908

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#### (2) INFORMATION FOR SEQ ID NO: 88:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 655 base pairs(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 88:

40 TGCACTGGTT CCTTCTCCCC AGCAAATACT GCCTTCTTGT TTTTCTCTGA TGTGGCAGGT 60 GACTACAAAA TCCGCCTTGG TATTCTTCAA ATGCATATAT ATTCCTTTCT TGTCAGCTCC 120 45 CTCTCTTCCT AGATTAGAAA ACTGCCTCAT TTTCTGCTCA CTGGATGTGC AGTCCCAGCT 180 240 TGTCTTCCTC TCCTCCCCCC CTGTTGCAGG TGTTCTTTTT TTTTTTCTTC TCTCCCCACT GGGCAGCAAA AGTTGTTCCA CAGTGGAAAW TTAGGCATCC TCAAGTTTCY TCCCAGCTTC 300 50 TGCTGTGTTT TCTTAGAGTA AATTGCCAAT TTCTGTTTTT ACAGGAAATC CTTTTTTAAA 360 AATGGAATCA GTGTGGTCCC CATCTACTCT GCAAAAATTG CATTTTTCTC TATTTTCAAA 420 55 480 CTTGGGCATT GCTWGATATG TGAAATGGGT TTATGAAAAA TAATAAAATC ATAACGCTAT 540 TTGTTTGACT TTCAATTTCA TGGGAATTTT TCTCAGCTAA ACTCTAAATG GTGATTARGC 600 60

120

# AAAAAAAA AAAAAAACY GRAGGGGGC CCGGTACCAA TTCGCCCTAT AATGA 655

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(2) INFORMATION FOR SEQ ID NO: 89:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1102 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 89:

TTTTTTTTT ACCATTTAAA ATAAAATGAA AGTGACCTTC TGTTTATAAA AATCTTTGTC
TGCATCTCTG CTTATTTCCT TAGAAGAGAT TCCAAGAAGC GGTGAGTGAT TTCACGGCAG

20 CAGAGGGTTG GGACATATTA CGGGCGCGGA TCCCTCTTGG AGTGAGATGA CTCTCCGGAG 180

AGATTTAGTC GTCACCCTCG CGTGTGAGGC TGCGTCACAC CCCAGGGATG TGTCTATCAA 240

GATGGAAGAT CTTTTACACG CTCTTGATTT TGTTTGSCTY TTTTTCTATT ACTAGTGAGA 300

25
AKGAAACTIT TTATATGATT ATTATCCATC ATAATCCAAC ACAAATTACT GCTTCATGTT 360

CTTTTACTTT CCTGTGAAGG TTTTAGTGCC TTTTAAAAAT TGCTATATAT TAAGCTTGTT 420

30 AATACTTCCA TGCTGTATTT GTGGSCATCA RTTTCCCCGG GNACAGGCNT GCACATTTTG 480

CCTTCACACG CTGGGTGGTT TTTCATTTTC AMITCTATTT CTCGTTCTTC TATCGTTTTA 540

TGTTCAGACG GGTTTCTCCG TGTAGAAAGC AGTTTATGAA GATTTACTTT CGACAGTCTT 600

CTCTCTACTT TCTACAGTGA ATTCTCTGAT GTGTCTGGGA GTTTGGGGGT CTGGGTAAGA 660

RTCCTCCTCT CACCCTATTC TCTATTACGA TCCACAGCCT CATGCTTTAT GARATTGGTG 720

GCCGGGARCG GGGGAGATTT GCGGATCCCC CAAGCCAGAC TTTATCCCCC TATCCCTGCC 780

TCTGGATCCC ACGTACAGGC CTGGGAACTC CCTGTGGGTA GGGGCCAATG GTCTCGCACT 840

CTCACCTGTA CCCCAGGGCT GGCACAGGAT GGTCAAGGAG AGAGGCTGCC CAAGCGCATC 900

CYTCTGGTGT CCCCCTGACA CGCCTCCAAA GTGAGCAGGT AGGTTTCAAC AGCCCCACGT 960

TGCAGGTGGG AGATGAAGCT CAGGGTGGAG ACCAGTATCT CACAGTTCTC TTTGCATGGC 1020

CGGGTACTTG TTAGTCAACT GATCAAGTGA AAATTCTAGC CCCAGAGGCA GGAGAATCCG 1080

GAACAAAATT AAACCAGCCA GG 1102

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(2) INFORMATION FOR SEQ ID NO: 90:

(i) SEQUENCE CHARACTERISTICS:

60 (A) LENGTH: 1533 base pairs

WO 98/54963 PCT/US98/11422

343

(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 90: GGCACGAGCC GNCACGGGCA GCGCCCCATA GCGCCAGGGA CCCCCTGGCA GCGGGAGCCG 60 CGGGTCGAGG TTATGGATCC AGCGGGGGG CCCCGGGGGG TGCTCCCGCG GCCCTGCCGG 120 10 TGNCTGGTGC TGCTGAACCC GCGCGGCGGC AAGGGCAAGG CCTTGCAGCT CTTCCGGAGT 180 CACGTGCAGC CCCTTTTGGC TGAGGCTGAA ATCTCCTTCA CGCTGATGCT CACTGAGCGG 240 15 CGGAACCACG CGCGGGARCT GGTGCGGTCG GAGGAGCTGG GCCGCTGGRA CGCTCTGGTG 300 GTCATGTYTG GAGACGGGCT GATGCACGAG GTGGTGAACG GGCTTCATGG AGCGGCCTGA 360 CTGGGAGACC GCCATCCAGA AGCCCCTGTG TAGCCTCCCA GCAGGCTCTG GCAACGCSCT 420 20 GGCAGCTTCC TTRAACCATT ATGCTGGCTA TRAGCAGGTC ACCAATGAAG ACCTCCTGAC 480 CAACTGCACG CTATTGCTGT GCCGCCGGCT GCTGTCACCC ATGAACCTGC TGTCTCTGCA 540 25 CACGGCTTCG GGGCTGCGCC TCTTCTCTGT GCTCAGCCTG GCCTGGGGCT TCATTGCTGA 600 TGTGGACCTA GAGAGTGAGA AGTATCGGCG TCTGGGGGAG ATGCGCTTCA CTCTGGGCAC 660 CTTCCTGCGT CTGGCAGCCC TGCGCACCTA CCGCGGCCGA CTGGCCTACC TCCCTGTAGG 720 30 AAGAGTGGGT TCCAAGACAC CTGCCTCCCC CGTTGTGGTC CAGCAGGGCC CGGTAGATGC 780 ACACCTTGTG CCACTGGAGG AGCCAGTGCC CTCTCACTGG ACAGTGGTGC CCGACGAGGA 840 35 CTTTGTGCTA GTCCTGGCAC TGCTGCACTC GCACCTGGGC AGTGAGATGT TTGCTGCACC 900 CATGGGCCGC TGTGCAGCTG GCGTCATGCA TCTGTTCTAC GTGCGGGCGG GAGTGTCTCG 960 TGCCATGCTG CTGCGCCTCT TCCTGGCCAT GGAGAAGGGC AGGCATATGG AGTATGAATG 1020 40 CCCCTACTTG GTATATGTGC CCGTGGTCGC CTTCCGCTTG GAGCCCAAGG ATGGGAAAGG 1080 TGTGTTTGCA GTGGATGGGG AATTGATGGT TAGCGAGGCC GTGCAGGGCC AGGTGCACCC 1140 45 AAACTACTTC TGGATGGTCA GCGGTTGCGT GGAGCCCCCG CCCAGCTGGA AGCCCCAGCA 1200 GATGCCACCG CCAGAAGAGC CCTTATGACC CCTGGGCCGC GCTGTGCCTT AGTGTCTACT 1260 TGCAGGACCC TTCCTCCTTC CCTAGGGCTG CAGGGCCTGT CCACAGCTCC TGTGGGGGTG 1320 50 GAGGAGACTC CTCTGGAGAA GGGTGAGAAG GTGGAGGCTA TGCTTTGGGG GGACAGGCCA 1380 GAATGAAGTC CTGGGTCAGG AGCCCAGCTG GCTGGGCCCA GCTGCCTATG TAAGGCCTTC 1440 55 TAGTTTGTTC TGAGACCCCC ACCCCACGAA CCAAATCCAA ATAAAGTGAC ATTCCCAAAA 1500 AAAAAAAAA AAAAAAAAA ANCCCGNGGG GGG 1533

WO 98/54963 PCT/US98/11422

344

	(2) INFORMATION FOR SEQ ID NO: 91:	
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 575 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: double</li> <li>(D) TOPOLOGY: linear</li> </ul>	-
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 91:	
	ATCCTCTGGA ATCTAGGTGG AAGCCACCAA GCCTTCTTCA CACTTGCGTT CTGAGCATCT	6
15	GCAGACTTAA CCCCATGTGG CAATCACCAA GGCTTATGGC TTGTGTCCTC CAGAACTGTG	12
10	GCCAGAGCTG TACCTGGGCC CCTTTGAGCT GAGGCTGAAG CCAGAGTCTG AAGCTCAGCA	18
	GGGCAGTARG GCCCTGGGCC TGGCCCCTGA AACCATTCTT TTCTCCTAAG CCTCTGGGCC	. 24
20	TTTGATGGGA RGGGCTGTCC TCAAGATTTT TGAAATGCCT TTGGAGGGTT TTTGCCTTGT	30
	CTTGGATATT GGCTTCCTTT TAGTTATGCT CATCTCTCTA GCAAGTGAAT GTTTCACAAC	36
25	CTGCTTGGAT TCTTTCTCTA CCACAGARCC AGGCTGCAAA TTTTACAAAC TTTTACACTC	42
	TGTTTCCCTT TTAAATATAA ATTTCAATGT TAAGTCACTT CTTTGCTCCC ATATCTGATT	48
	TAGGTTGCTG GAAGTAGCCA AGTCACCTCT TGAATGCTTT GCTGCTTAGA AATTTCCTCT	54
30	ACTAGGTAGC CTGGGTCATC ACACTTAAGT TCAAA	57
35	(2) INFORMATION FOR SEQ ID NO: 92:	
40	(i) SEQUENCE CHARAÇTERISTICS:  (A) LENGTH: 639 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	,
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 92:	
45	TCCTTTCATC TTAAGCACCA CCCGACAGGG CAGGTACTAT TACCATCTCC GTTTGACAGA	6
	TNAGGAACCT GGCACAGGAA GCATTTAAGT GGATTCCCCA GGATCGCCCC ACTGTCAGGA	12
50	GCAGANTCAG AATGGGCCTC AGCATCAGGC TCCCAATCCT GGCTTCTAAC TGCTGCGCTC	18
	TSCCCTTCYC TCWCCCCACC TCCCCACTCC AGTGCCTTTG GTCATGCCAC TGCAGCTTTC	24
	AGGCCAATAC TGGATTAGCC TCTTAGTGTT CTTGTCCCTG CAGCCATTTC CCCAGGCAGC	30
55	AATTCCATGT GCCCTCACTG ATGTAGGTGG CTCTTGTGTC ATTTGTCACA TCCTATTGAA	36
•	TTGTTTATGC ATCTTGTTCA CACTCACAGC ACCCTCCCTC TCACACGTCC TCCTTATAAA	42
•	AATGTCCCTC AGTGTCTGCT ATGAGCCAGG TGCAGACTTA AGTGACAGGG CTGCTACGGG	48

	•.	
	AAATAAAAAA TTAACAAGGA GCACCTGCCT CTTAATGCAC AGTAACAAAC TATGTTAAGT	540
	GTCAGGAAGG AAAGGTTAAG GATGCCAGGA AGGCTTTTAA TAAATAACCT GACTTAGATG	600
5	GGCAGGTGGT GCTGARGATT AAGAACGTGT TCTTCTCGA	639
10		
10	(2) INFORMATION FOR SEQ ID NO: 93:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 744 base pairs	
15	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
13	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:	
20	GAATTCGGCA CGAGAGTGGC TGGAGTCTGG CTGCAGAGGG AAGACATCAG CAGGGAGGGA	60
	GCCAGGGCCT GTCACATCTT TCCTCTGGCC ATTGTCCTGG TCTTTGTAAG CCCAGAATCT	120
25	CCCCTTCCCT GAAGGGAGGC CAGCACCCCA GGAGGGCAGC AGGTGTGCTG TGAGGGTTGG	180
	AGTAGTGTGA GAGGTCAGGG TACACTAGAA TGGCCATGGA CACCATGTGG GGGTGCTCTG	240
	GGCTGGGCCA CAGAACAGTG TCCTTCCTGC TGCTCCTCCC CTGCAGCTTC CCCCGACCTT	300
30	GTNGTTTATT TGGTTTGATA CCAATCAGCA GACCCTGCAA GGTGGAAGCT CCCAGGCTCT	360
	CAGTCCCACS ACTCTCATGT GCCAGTCACC CNTACTGTAA CTGCCCAATG AGTACTTCTT	420
35	GCCCACTGCC AAGATAGAGC CAGTTTACCA AGACAGGGGA ATTGCAGTAG AGAAAGAGTT	480
	GAATATACAT AGAGCCAGCT AAATGGGAGA GTGGAGTTTT CTTATTACTT AAATCAGCCT	540
	CCCYTAAAAT TCAGAGGTGA GAATTTTTCA AGGACAGTTT GGTGGSCAGG CCTAGGGAAT	600
40	GGATGCTGCT GATTGGCTAG GGATGCAATC ATAGGGGTGT AGAAAAGTWC CTTGTGCACT	660
	GAGTCCACTT TTGGTGAGAG CTACCAAGGA GCTGCTGGTC TGCTGGTCCC GGTAGAGCCA	720
45	TCTGGTGTCA GGAATGCAAA AGTG	744
	(2) INFORMATION FOR SEQ ID NO: 94:	
50	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 526 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 94:	
60	GCAGGGGAAT TCGGCCACGG AGGGGTTTCA ACAGGGCCCG TGGGGTGAGG TGCARACACA	60

PCT/US98/11422

	AAGCCCATAA GTGCTGGCCT GTTGGGACAA ATGAGAGAAA TCCCATAGGG TGGTGATGAC	120
	AGCGCAYTCA GCCATCYTAY TCCTGGGGAA AATGAAACTT GTGCTCCTAT CAAATGCTCA	180
5	GTTGTAAAAC TGGAAAAAAA TTTTAGAAGA CATCTTGTCC AGCATCTGTG TTTATGTCTA	240
	TAAAATGTAG AAAACTAAAG CACAGAGATG TTAAATGTTT TGTCCAAGGT CCAACAGCTG	300
10	GTTAGCARGC TTGGTCTGGT GACCTTTCTA CTGAACCACA GTGCCGCTGG GGGAAGTCCT	360
10	CAGCACAGAT GGCTGCTGCT ATAGCTGGGG TATGGGCAGT ATTAGTAGTT AACCAGTCAA	420
	CCCAAGTTCC CATAGTCTAG GTTCTGCTTC AGCTGGAGGT TAGGGAAAAA CACAAGAAAA	480
15	TCCCTTACCA CTCTACCAGT GCTGGGGGAT GTACTAAGAG ATCCCC	526
20	(2) INFORMATION FOR SEQ ID NO: 95:	
25	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 426 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li><li>(D) TOPOLOGY: linear</li></ul>	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 95:	
30 .	GGCACAGGGC AGGAGAGACT TGGTCCATGG GGAGAAGCCT GCAGTATAGA TGGGACCTCC	60
	AGGAGCCCAA GTAGCATAGA CCCTGCTGAT CCGGGGCCAT TGAGCCAGAG GATTTGGGCT	120
25	GAATGTCCCC AGAGACAAAA GGGAAAGGTA GATCCTTTCC CTTAAAGATG AAAGCCATCG	180
35	CCCGGGCTTG CTTATTGCTC TCTCTCCTGG TCCTTCCACA TGTTGTTTCT GAACATTTGT	240
	TCTGGCATCA CAATCCCCGT CATCCTGTCA TCTGGCCCTT CCCACCTTTC CACCTTATCT	300
40	CTTGCAGTGT CTCCGCGTCG ACCTGGCACC TGGGTGAARG CTTGCTCTTG CTGGTGCCCA	360
	TAGCCCCCAG TGTATGGTCT TGAMCTCCCC AGCCATATGG ARACCCACCT CAGGAGGGCC	420
45	CCTCGA	426
	·	
50	(2) INFORMATION FOR SEQ ID NO: 96:	,
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 844 base pairs  (B) TYPE: nucleic acid	
55	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 96:	
60	GCCACAGCGG CACGAGATAG GAAGCTTGGC AGGGGCAGCT CCCCCAGTGC GCATTGCCCT	60

	GTAACTCGAG	CGCCTGGGAG	TGGGGAGAGG	CTTGGAAATG	GAGCAGGGTG	GTGGACCTCG	12
	TCTTCTCCTG	CTCATCCCAG	GCCTCCTCCA	TAACACCTAC	CTAGCACGGC	CTGGGGACTT	18
5	CCCAGCCCAA	GGAACAACTG	AGAATACTGA	GTGCCAGGGT	AGCCCTAGCC	CCATTTCACA	24
	CCTGGGCAAA	GTGAGGTCAC	TGGATTCAAA	CACTCAGATT	TAAACCTCCT	CTGTGTCTGC	300
10	AGCACCTGTA	TATAACTGCC	AGCCTCTGCT	GCCCCTCTCC	AAAAAGTCTC	TGCCCTTGTC	360
	TTTGGCACCT	GTCTCTGTCC	TCCCCATTCT	CTGCTCCTCC	TTTCTCCAAC	TCAGANTCAC	420
	CCTGTTAGTT	CAGCAAATGT	TCATCGAGCT	CCATAATGTA	GCAGGACAGG	NCTGTCTAAC	480
15	AGATTCTGGN	CTTGCAAGGG	TGAGACAAGT	ACTCTCCATC	TTTCTCTCAT	CTTCACAGAT	540
	GGTCTGCTCA	ACAACTTTGC	ACTGAATTGT	AAATAATTGA	TACTGCATAA	AACATTGATG	600
20	TTCTTTAAGG	GTAGTCCAGC	AAGGTGGCAA	GTCTTATAAT	GATAACTGCT	CAAGGATCTC	660
	TCAGTGAAGC	ATTTGGGGST	GCTAGCTCTG	CCTATGGGTG	AGGTCAGCTA	TCTCACGCCA	720
	TCTACTTCCA	CNTGCCCCCC	CATGCCAGGC	TCACCCTGAG	CTGAGATGCC	TGAGCAGGTG	780
25	GCAGAAAGGA	GCCACCTGGT	TTATGCTTCG	GGACCACAAA	CTCCTCTATC	CAGANGACAG	840
	TTTT						844

35

# (2) INFORMATION FOR SEQ ID NO: 97:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1985 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 97:

AGCCCTGCTG AAGTACAGGT TCTTCTATCA GTTTCTGTTG GGCAATGAAC GAGCAACAGC 60 AAAGGAGATC AGGGATGAAT ATGTGGAGAC GCTGAGCAAG ATTTACCTGT CTTACTACCG 120 45 CTCTTACCTG GGGCGCTCA TGAAGGTGCA GTATGAGGAA GTCGCTGAGA AAGATGATCT 180 AATGGGTGTG GAAGATACAG CAAAGAAAGG ATTCTYCTCA AAGCCATCGC TCCGCAGCAG 240 50 GAACACCATT TTCACCCTAG GAACCCGCGG CTCTGTCATC TCCCCCACTG AACTTGAGGC 300 CCCCATCCTG GTGCCTCACA CAGCGCAGCG GNAGAGCAGA GGTATCCATT TGAGGCCCTC 360 TTCCGCAGCC AGCACTACGS CCTCCTAGAC AATTCCTGCC GCGAATACCT TTTCATCTGT 420 55 GAATTTTTTG TTGTGTCTGG CCCAGYTGCA CACGACCTGT TCCATGCTGT CATGGGCCGT 480 ACACTCAGCA TGACCCTGAA ACACCTGGAT TCTTATCTAG CTGACTGCTA CGATGCCATT 540 60 GCTGTTTTTC TCTGTATCCA CATTGTTCTC CGGTTCCGTA ACATTGCAGC AAAGAGGGAT 600

	GTTCCTGCCC	TGGACAGGTA	CTGGGGAACA	GGTGCTTGCC	TTGCTATGGC	CACGGTTTGA	660
5	ACTGATCCTG	GAGATGAATG	TTCAGAGCGT	CCGAAGCACT	GACCCCCAGC	GCCTAGGGGG	720
J	GTTGGATACT	CGGCCCCACT	ATATCACACG	CCGCTATGCA	GAGTTCTCCT	CCGCTCTTGT	780
	CAGTATCAAC	CAGACAATTC	CTAATGAACG	GACCATGCAA	TTGCTGGGAC	AGCTGCAGGT	. 840
10	GGAGGTGGAG	AATTTTGTCC	TCCGAGTGGC	AGCTGAGTTC	TCCTCAAGGA	AGGAGCAGCT	900
	TGTGTTTCTG	ATCAACAACT	ATGACATGAT	GCTGGGTGTG	CTGATGGAGC	GGGCTGCAGA	960
15	TGACAGCAAA	GAGGTTGAGA	GCTTCCAGCA	GCTGCTCAAT	GCTCGGACAC	AGGAATTCAT	1020
	TGAAGAGTTG	CTGTCTCCCC	CTTTTGGGGG	TTTAGTGGCA	TTTGTGAAGG	AGGCTGAGGC	1080
	TTTGATTGAG	CGTGGACAGG	CTGAGCGACT	TCGAGGGGAA	GAAGCCCGGG	TAACTCAGCT	1140
20	GATCCGTGGC	TTTGGTAGTT	CCTGGAAATC	ATCAGTGGAA	TCTCTGAGTC	AGGATGTAAT	1200
	GCGGAGTTTC	ACCAACTTCA	GAAATGGCAC	CAGTATCATT	CAGGGAGCGC	TGACCCAGCT	1260
25	GATCCAGCTC	TATCATCGCT	TCCACCGGGT	GCTGTCCCAG	CCGCAGCTCC	GAGCCCTCCC	1320
	TGCCCGGGCT	GAGCTCATCA	ACATTCACCA	CCTTATGGTG	GAGCTCAAGA	AGCATAAGCC	1380
	CAACTTCTGA	TGTGCCAGAA	ACCGCCCTGA	GATCTGCCGG	TCATCTCCAT	GGACTTCTGC	1440
30	ACCCCATTCC	ATACCCTTCT	TCACCTGGGG	TACCCCTTCC	AGTTTTCCCC	TTGCTTCCCA	1500
	GGCCCTTGAC	ATGGCTTACC	TGCCTTCACT	CCCAGCACCT	TGCCCAACAG	GATAAGCTGG	1560
35	ATCCCCTTGG	CCTTCTGAAT	ATCCCAGTGT	CTTCAGGTTT	CCCAAGACCA	CTTCCCTGTG	1620
	GGCTTCCAAA	ATGGCCTTTA	TCATTTCTCC	AGTCTGTCAC	CCTCCTTTCC	TGCTCCCATA	1680
	CACCCAAGGC	TTGTTTCTTC	CCCTGTAAAA	ACCACTGCCT	CAATCTCTGG	TTCACTCAAC	1740
40	TAGTCACCAT	GTCCTGAGGC	ATGAAGCCTC	CTCAGCTCTT	GGAATTGCTG	GCAAGGGGTG	1800
	ACTGCCTCTG	AGTCATTGTG	TTTTTCAAAG	TGATTTCTTT	TCTGTAGCTT	TTTGACCTAA	1860
15	GATCTCAGCA	ATTTGAACAC	TAACCTCTCC	CCTCCTGGCT	CAAGAATTAC	TCCGAAGTCA	1920
	GTCTGCAGAA	AATAAATATT	TAGTATGACA	TGAAAAAAA	АААААААА	АААААААА	1980
	AAAAA		•				1989

# (2) INFORMATION FOR SEQ ID NO: 98:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1416 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

#### (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 98:

	AIAIGAAGGG	MAGMITIG	AIIAIGIIII	CICANIIGNI	GICARIGANG	GIOGACCAIC	00
5	ATATAAATTG	CCATATAATA	CCAGTGATGA	CCCTTGGTTA	ACTGCATACA	ACTTCTTACA	120
	GAAGAATGAT	TTGAATCCTA	TGTTTCTGGA	TCAAGTAGCT	AAATTTATTA	TTGATAACAC	. 180
10	AAAAGGTCAA	ATGTTGGGAC	TTGGGAATCC	CAGCTTTTCA	GATCCATTTA	CAGGTGGTGG	240
10	TCGGTATGTT	CCGGGCTCTT	CGGGATCTTC	TAACACACTA	CCCACAGCAG	ATCCTTTTAC	300
	AGGTGCTGGT	CGITATGTAC	CAGGTTCTGC	AAGTATGGGA	ACTACCATGG	CCGGAGTTGA	360
15	TCCATTTACA	GGGAATAGTG	CCTÁCCGATC	AGCTGCATCT	AAAACAATGA	ATATTTATTT	420
	CCCTAAAAAA	GAGGCTGTCA	CATTTGACCA	AGCAAACCCT	ACACAAATAT	TAGGTAAACT	480
20	GAAGGAACTT	AATGGAACTG	CACCTGAAGA	GAAGAAGTTA	ACTGAGGATG	ACTTGATACT	540
	TCTTGAGAAG	ATACTGTCTC	TAATATGTAA	TAGTTCTTCA	GAAAAACCCA	CAGTCCAGCA	600
	ACTTCAGATT	TTGTGGAAAG	CTATTAACTG	TCCTGAAGAT	ATTGTCTTTC	CTGCACTTGA	660
25	CATTCTTCGG	TTGTCAATTA	AACACCCCAG	TGTGAATGAG	AACTTCTGCA	ATGAAAAGGA	720
	AGGGGCTCAG	TTCAGCAGTC	ATCTTATCAA	TCTTCTGAAC	CCTAAAGGAA	AGCCAGCAAA	780
30	CCAGCTGCTT	GCTCTCAGGA	CTTTTTGCAA	TIGITTIGIT	GGCCAGGCAG	GACAAAAACT	840
	CATGATGTCC	CAGAGGGAAT	CACTGATGTC	CCATGCAATA	GAACTGAAAT	CAGGGAGCAA	900
	TAAGAACATT	CACATTGCTC	TGGCTACATT	GGCCCTGAAC	TATTCTGTTT	GTTTTCATAA	960
35	AGACCATAAC	ATTGAAGGGA	AAGCCCAATG	TTTGTCACTA	ATTAGCACAA	TCTTGGAAGT	1020
	AGTACAAGAC	CTAGAAGCCA	CTTTTAGACT	TCTTGTGGCT	CTTGGAACAC	TTATCAGTGA	1080
40	TGATTCAAAT	GCTGTACAAT	TAGCCAAGTC	TTTAGGTGTT	GATTCTCAAA	TAAAAAAGTA	1140
	TTCCTCAGTA	TCAGAACCAG	CTAAAGTAAG	TGAATGCTGT	AGATTTATCC	TAAATTTGCT	1200
	GTAGCAGTGG	GGAAGAGGGA	CGGATATTTT	TAATTGATTA	GTGTTTTTT	CCTCACATTT	1260
45	GACATGACTG	ATAACAGATA	АТТААААААА	GAGAATACGG	TGGATTAAGT	AAAATTTTAC	1320
	ATCTTGTAAA	GTGGTGGGGA	GGGGAAACAG	AAATAAAATT	TTTGCACTGC	TGAAAAAAA	1380
50	АААААААА	AAAAGGAAAC	TCGAGGGGG	ccccc			1416

# (2) INFORMATION FOR SEQ ID NO: 99:

55

60

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1935 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 99:

5	NTCTACCCTA	ATCAAGATGG	GGACATACTT	CGCGACCAGG	TTCTTCATGA	ACATATCCAG	60
J	AGATTGTCTA	AAGTAGTGAC	TGCAAATCAC	AGAGCTCTTC	AGATACCAGA	GGTTTATCTT	120
	CGAGAAGCAC	CATGGCCATC	TGCACAATCA	GAAATCAGGA	CAATAAGTGC	TTATAAAACC	180
10	CCCCGGGACA	AAGTGCAGTG	CATCCTGAGA	ATGIGCTCTA	CGATTATGAA	CCTCCTGAGC	240
	CTGGCCAATG	AGGACTCTGT	CCCTGGAGCG	GATGACTTTG	TTCCTGTGTT	GGTGTTTGTG	300
15	TTGATAAAGG	CAAATCCACC	CTGTTTGCTG	TCTACTGTGC	AGTATATCAG	TAGCTTTTAT	360
13	GCTAGCTGTC	TGTCTGGAGA	GGAGTCCTAT	TGGTGGATGC	AGTTCACAGC	AGCAGTAGAA	420
	TTCATTAAAA	CCATCGATGA	CCGAAAGTGA	CCAAGACCAA	GGCCCACCAA	GGCAGCAGAC	480
20	TGTTAATCAG	ACAAACAGAT	CTCTGAGAAG	GTGCATCAGC	TGCTTTGAAG	GCTGAAGATT	- 540
	GTTTTGTATG	ATACTGCACA	GCATCAGGCA	TTTTAAAGCA	GATCTTTACT	AAACAGGTTA	600
25	ATGAGCTAAC	AAGCAGGTTC	TCTCGTCTTT	GGGCTCTTTC	CTTTCTGAGT	TGCATATTCT	660
25	ATTTTCTTGT	CCCCAAGTAG	AGACTAGTAC	TACAAAAAGG	GACCACATTT	TTCAAGTATT	720
	TCTAAGTATA	AAAAACAAAA	CAAAAATCTC	TTAGGAAATG	TCTAGACCTC	CATTCTTGGA	780
30	TTCCCTTTCT	TTCCTTTTAT	TTTAAAAAAG	AACAGTACCC	CTCTTTTAAG	ATGCTGTCTT	840
	ACATTAATGA	GCATCTAATG	GAAAGAAGGT	ATGAGTTGCA	CTGAGGATTA	GAATAGTGGT	900
25	GCGTTAGTGG	CATTATCTAT	AAATACACTC	ACCTAAATTG	AAAGCTAAGA	AGGAAATGTA	960
35	AATATAATAT	ATATTTATAT	TTGATGTAAT	ATGGACATCT	GCAGATTCTA	ATAAACAAGG	1020
	ACTATIGCTG	ATAGTAGGCT	GTGACATACT	GTCTTGTGAA	ATGGTTTCCT	TGACAAAATT	1080
40	TAAGCTGAGC	TTAAAAGCAA	AAAAACAAAA	AGTACACAGA	AATATTTATT	AAAATGTAAT	1140
	ACAGTTTATT	GAACTTTCTA	GGTATGGAGT	TTGATGGACA	GGGCTGCCTY	TAATGAGTGT	1200
45	GAAGGTCACT	AAGTCACTTA	GACATCTCAC	CGTGGAAGTT	TGTGAGCCTG	CATTAGGAGA	1260
43.	TAGACTGATT	ACCATACATG	ACATAAAAAG	GAACAGTGGA	TAGCTCATAC	TTTATGGTGG	1320
	TTCTTCTCCT	CCGAAATAAT	ATACTGCAGA	AATCCCAGAC	AGAGCTCCTT	ACAAACCTTT	1380
50	AATTGTAATA	TATTTTTGAT	GATTATTCAC	ATTGAATGCA	CAGACCAAGA	ATTCAGTGAA	1440
	TGTCATTTTT	тааааааста	ATTTGTATTG	TCTGCTCTAG	TGATACAAGT	TTTACTAGTG	1500
55	ATAAACTATT	TTAATCAACC	ATACTATTCT	TATGGAAAAA	AATATCTATT	TTGGCAGGTT	1560
55	TCTGTGCCTT	TATTTCCCTC	TTCTGAAAAA	AAGTCTGTGT	TTTCATAGTT	TGGTTTGCAT	1620
	TGTATATCAA	тааттаатса	GGAATGGGTT	TTGGTGCCTG	AAAAATTGGC	CATGGAGGCA	1680
60	CACCAAAGCT	TCAAGCACAA	GTCTTGTACA	TGGGCCATCA	CTGTCTGGTT	TCACTTCGTG	1740

	TGITTCCTAA ACACATTTAG CTGCTTTTTT AACAAACTCA GCCCCATACT TGAGTCCCTT	1800
5	GTTGTTGGGA GCATTTCCAG GCATCTTTTA AGGGAACTGT GACAAACAGC CTCGGGCAGA	1860
, •	TGAACACGGA GCCTCTCTGT TGTCTGTCTC TGAGATCTTT GTGTCTGGGA ATGCCTAAAG	1920
	NITTTGNITT TITTT	1935
10		
	(2) INFORMATION FOR SEQ ID NO: 100:	
15		
13	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 599 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 100:	
	GAATTCGGCA CGAGCGTCCA CGCAGCCGCC GGCCGGCCAG CACCCAGGGC CCTGCATGCC	60
25	AGGTCGTTGG AGGTGGCAGC GAGACATGCA CCCGGCCCGG	120
	CCTCATCCTG ATGGGCACTG AACTCACTCA AGACTCCGCT GCCCCCGACT CCCTGCTGAG	180
30	AAGTTCAAAG GGCAGCACGA GGGGGTCTTT GGCTGCTATT GTCATCTGGA GGGGGAAGAG	240
-	TGAGAGCCGG ATAGCCAAGA CCCCAGGCAT TTTCAGAGGT GGCGGGACCT TAGTCCTACC	300
	CCCAACACA ACCCCTGAGT GGCTCATCCT CCCTTTGGGC ATAACGCTGC CCTTGGGGGC	360
35	TCCAGAAACA GGCGGTGGGG ATTGTGCCGC TGAGACCTGG AAGGGCAGCC AGCGTGCCGG	420
	CCAGCTGTGT GCATTGCTGG CTTAATATGC AGGCCTTGGG GGGCTGTGGC CACATGCCCG	480
40	GCAGGAGGTG AGTGAGGAGC CCTGTGGCGT GCTGGTGTGG GGATCGTGGG CATTTCAAAC	540
	GGGCTTGTCG TACCCTGAAC AATGTATCAA TAGAGAAAAA AAAAAAAAAA	599
45		
	(2) INFORMATION FOR SEQ ID NO: 101.	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 784 base pairs	•
50	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 101:	
<i></i>	GAATTCGCCA CAGAAAAAAA AGAGAGACTG GGTCTTACTG TGTTGCCCAG ACTTGTCTTG	60
	AACTCCTGCC TCAGCCTCTC AAGTACTTGG GATTATAGGC CAAGAAGCCA CCATGCCTAG	120
60	CTTCTTCCTG TCATTGATCC AGACTAATAC TCTGGGGTCA GCCTCATTTC TTCTCTTTCT	180

	CACTITICAC ATCCACTIGT CACCAAATCK RGTTCATTCT GCATCCTAAG TAAGTCCTTT	24
5	GATTCCTCCA GTTGTTCATT AGTAATGTCT CAARTGTAAT TTTTTCTAGT AGTTTTCAGC	30
	CTGTCTTTCC KGCCTTCAGT CTTAACTTCT CCAGTACATA KGCCACATTG TTGTCAGCAK	360
÷	GATCAWATTT TATTTAAAAA TACTTTACAW AKGTTTATKG CCAAATATTA GRAAATACAG	420
10	ATTCATGGAA AGAAAAATCA CTGTCCCAAG GAGGTCACTG GCATGGTGAG GTTAAGGGGT	480
	GATTTTAATT TITAAAAATG TATATTTTTT CCTGTGTAGA GTAGTAACAC CCTTGAAAAC	540
15	ACAWTCCCTT GTAAAGTCTC TAATTCTGTA CTCCGCATCT AGSTGRTCTC TTCTTTCTCA	600
	GATATTTTAC AATTTCATTT ATCACCACCT TTCTCTAGCC TTTACCCGTC TCTTCAATAT	660
	TWACATATGC AGAAGTTTCT CCTAACAAAC ACCTGCCTCT GCCTCAGTTC TGCTACCACC	720
20	CTGTTGCTTT CTTTCCCTTC ACAATCAAAT TTAAGAGTGT CAAAAAAAAA AAAAAAAAAA	780
	TCGA	784
25		
	(2) INFORMATION FOR SEQ ID NO: 102:	
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1035 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 102:	
	AGAGGCCTGG CTGCGTTGCC CTATCTCCGT CTCCGCCACC CACTTAGCGT TTTAGGCATC	60
	AATTACCAGC AGTTTCTCCG CCACTATCTG GAAAATTACC CGATTGCTCC CGGCAGAATA	
40		120
	CAAGAGCTTG AAGAACGCCG CAGTTGCGTG GAAGCCTGCA GAGCAAGGGA AGCAGCGTTT	180
45	GATGCCGAAT ATCAGCGAAA TCCTCACAGG GTGGACCTCG ATATTTTAAC CTTTACGATA	240
45	GCTCTGACTG CCTCTGAAGT TATCAACCCT CTGATAGAAG AACTTGGTTG CGATAAGTTT	300
	ATCAATAGAG AATAGTTAGG TGGTGACACT ACTTCAAGAG AACCTCTGCA TTCCAGTCAT	360
50	ACCAATCCTG CAACTTGATT TTCAGAAGTC AAGAGTATAT CGCGATAAGA CAGTGCACAG	420
	GTGGAGGGGA AAAAAAGGGG GAGGGGGAAG CTTATCTTGA AAAAGCATCA CAGAAGTAGA	480
	AAAAAATGTC GAAAGCATTA TAACTGTAAC GTTCTTTGAG TTTGTGATTG ATCCACATTT	540
55	TTCCCCCTGC ATTATGGAAA ATGTCTCTCA GCATTGCTTT ATTACAAAGT AAAGGATGGT	600
	TTTATAAAAT TGAGACTGAT GAAACATCAA TACTAGAGCC CATGAGGATG AAAGAAATTA	660
60	TCAAATAGTG CTGAACAGAA TAAGATGTTA ACGCTGAGTT ATTAGGACTG GAAGGCTATG	720

	AAAAGAACTT	GAAATTGTCG	GAATATGTGC	TCTCTTCATG	TCATATTCAA	TAGAAGTTTC	780
	TAGTTTAAGA	TTGATTTTGT	GTTTTCTTAG	GCATTTCAAG	TGACAAGCAA	AGTAAATGTA	840
5	TATATTATGT	GATAAATCAT	GTTTTCAAGA	ACGTCAAATT	TCTGGACTTT	TITCTITCAA	900
	TTTTTAATTT	TTAAAGTTTT	TTTGGTATTA	AAAAATCYAT	TCACAAGCCA	AAAAATWIWI	960
10	WAAATWIWCM	GCGAAAAGCC	ааааааааа	AAAAMMAGGG	GGGCCGGGC	CCCATCCCCC	1020
	CAAGGGGGTC	CNGNT					1035

# (2) INFORMATION FOR SEQ ID NO: 103:

# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2218 base pairs 20

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103: 25

	AGGTATTAGG	CCCTTTTGTG	GGAGCCCCAT	GTTTTGTTTT	TCTGAGTTGG	TGGGGAGGGA	60
	SGGAGGGGGA	GGGCTGAATT	GTTTTGCAGA	GGAAGATGGC	ATCTGTGCTT	TAAATTTCTC	120
30	ATTACTGGGT	TAGAAAACAA	AGAGGGAKTG	CCCTGCACAT	TTTCTTTTGT	GCTTTTAAAT	180
	GTTTCTTAAG	TTGGAACAGG	TTTCCTCGGG	CCTGTTTTGA	CTGATTGCTG	GAGTGCATTT	240
35	GATAGTTAAA	AATTACTAAT	TGGTTTTATT	TCCCTTCACA	CTCTGCCTCC	CCACTTCTCC	300
33	CCCCGTTACT	GAAAAATAAC	CATTITAGTG	TCAGGCTAGA	AATTGAATTG	CTGAGTTTTG	360
	TGTATCCTTT	AAATTAAAA	CCACAAGTGT	TTATTGTAGT	GGTTAAACTG	TAGCATCTCA	420
40	GCATCTGGGT	GGAAGCTGCC	TATATTTCTT	CCCAGTTTAA	CTGGGGACCA	TCTGTGAAAT	480
	TAATTTTCCA	TCCAGACAGC	TGCTGTGAGC	AAATGAACAT	AAATGCTCGC	TGGAAATTTA	540
45	CTAACCAGTT	TTTATATTGA	CCTGCAGTGT	AAAAAGCACA	TTTAATTATA	AACAATATAT	600
43	TCAAAATGGG	CAAATTTTAT	TTTCAAATGC	AGTGTAGAGC	TAGATTAAAA	GCAACTCTTT	660
	GCCACCTACT	CTGCCCTTTT	GGCAAAGTTA	CCTTGAACAA	AGAATCTTAA	GGGTTTATTA	720
50	AGAACTCTTT	ATTTTCTTCA	TACCCTGTTC	TCTGCAGTGC	TTTCTAACAG	CTTCTGGGTG	780
	CAGATTTTCT	TCGGCATCCT	TTTGCACTCA	GCTTATTACA	GGTAGGTAGT	GCTTAAGAAA	840
e e	AGTCATGGAG	GACTAAAGCC	TAAGTCCTTT	TCACTTTTCC	TCCATCTGAA	GGTAGGTGAG	900
55	TTCATCCTCT	TCATAGTAAT	GCTGTTTTAC	CAAGACTTTA	TAGCAGATGG	ACCCAGAAAG	960
	AATTTTCTGC	TATTGTGTTC	ACTACAACAG	GATAGGGACA	TCAGACAGCC	CCAGAAACCC	1020
60	CTTCCAGATC	TGATATGGGA	СТАТТААТТТ	TTATGCTGTT	AATTGGTATT	CATTCACAAT	1080

TOTAGGATCT ATACTCGAGG TITTGTTTTC CITTTANAAT TCTTTAGGGA GAGGAGG GTTCTGAAG GGTTCGAAA GTATGATTCA ATGTGCAACA TACAGGTAGG TCTTCAA AAGCTGAAAT ATATGCATGT AAAAACTTTG ACATCTTTT TITTAATTTT CCACTT.  AAGCTGAAAT ATATGCATGT AAAAACTTTG ACATCTTTTT TITTAATTTT CCACTT.  CTTAACTTTA CITCTCTTTT TGTCCCCCCC CCATCTTACA GAAGTTGAGG CCAAGGA ATGGTAGGCA CAGAAGAAAC ATGGCAAACT GCTCTGTGT TTCAAACCAA AGTGTTC CCAACCCCAA ATTTGTCTAA GCACTGGCCA GTCTGTTGTG GGCATTGTTT TCTACAA CCCACCCCAA ATTTGTCTAA GCACTGGCCA GTCTGTTGTG GGCATTGTTT TCTACAA CCTCTCTCGCA GCTCTTGAAA GCATCTGTTT GAGGGAAAGG TCTCTGGGCA AGCAAGT CTTTTGGATT GCTTGCTTCC CTTTTTCCAC CTGGGACATG GYAATCATAA AATAACA AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAACTCAAC AGAGTTT GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT TCTTGCTGGT CTCCAGGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT CTATTGAA TCTTGCTGTG CTCCAGGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAAAAA AAAAAAAAA AAAAAAAAA		GCAGTTGAAG	GGGGAAGGCT	CCACTGCATT	CTTTGGCTAA	GCCTGAATC	CTTGCTCATC	1140
AAGCTGAANT ATATGCATGT AAAAACTTTG ACATCTTTT TTTTAATTTT CCACTT.  CTTAACTTTA CTTCTCTTTT TGTCCCCCCC CCATCTTACA GAAGTTGAGG CCAAGGA ATGGTAGGCA CAGAAGAAAC ATGGCAAACT GCTCTGTGCT TTCAAACCAA AGTGTTC CCAACCCCAA ATTIGTCTAA GCACTGGCCA GTCTGTTGTG GGCATTGTTT TCTACAA CCAACCCCAA ATTIGTCTAA GCACTGGCCA GTCTGTTGTG GGCATTGTTT TCTACAA CCAACCCCCAA ATTIGTCTTA GCACTGGCCA GTCTGTTGTG GGCATTGTTT TCTACAA CCCCCCCAC GCTCTTGAAA GCACTGGCCA GTCTGTTGGG GGCACTGGTT TCTACAAAC CTCCTCGCCA GCTCTTGAAA GCACTGGTT GAGGGAAAGG TCCTCGGCA AGCACAGT AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAATCTAGC AGCACTT CTCTCTGCCT CCATGTCTGT CACTTATAAT TCAGGTTCT GTGTTGGCTT CAGAACAC GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCCTAACA ATTCGTT GCAGCTTCCCATGT TATTAAGCAA TGGGCTGCTT TTGGTTGGGA TCAGTGCTCT CTATTGAA TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG AAAAAAAAAAAAAAAAAAAAAAAAAAAAA	5	TGTAAGATCT	ATACTCGAGG	TTTTGTTTTC	CTTTTAAAAT	TCTTTAGGGA	GAGAGGGATG	1200
10 CTTAACTITA CITCTCTTT TGTCCCCCC CCATCTTACA GAAGTTGAGG CCAAGCC ATGGTAGGCA CAGAAGAAAC ATGGCAAACT GCTCTGTGCT TTCAAACCAA AGTGTTX CCAACCCCAA ATTTGTCTAA GCACTGGCCA GTCTGTTGT GGCATTGTTT TCTACAA  AATTCTGGGT TTTTTTCTTC TTTCTTTAAA CATAGAGGTA CCACCACAAG GGATGCC CTCTCTCGCA GCTCTTGAAA GCACTGTTT GAGGGAAAGG TCTCTGGGCA AGCAAGT  20 TATTTGGATT GCTTGCTTCC CTTTTTCCAC CTGGGACATT GYAATCATAA AATAACA AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAATCTAGC AGAGTTT CTCTTCTGCCT CCATGTCTGT CACTTATAAT TCAGGTTTG CTGTTGGCTT CAGAACA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT TCTTGCTGCT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGGTS TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGGTS  30 TGGACTTCTC ATCTAAAACG TTAGTGGCTT TTGCTTCGGA TCAGTGCTCT CTATTGA TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATS TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		GTTTCTGAGG	GGTTCTGAAA	GTATGATTCA	ATGTGCAACA	TACAGGTAGG	TCTTCAGCAT	1260
ATGGTAGGCA CAGAAGAACA ATGGCAAACT GCTCTGTGCT TTCAAACCAA AGTGTTC CCAACCCCAA ATTTGTCTAA GCACTGGCCA GTCTGTTGT GGCATTGTTT TCTACAA AATTCTGGGT TTTTTTCTTC TTTCTTTAAA CATAGAGGTA CCACCACAAG GGATGCC CTCTCTCGCA GCTCTTGAAA GCATCTGTTT GAGGGAAAGG TCTCTGGGCA AGCAACT  CTCTCTCGCA GCTCTTGAAA GCATCTGTTT GAGGGAAAGG TCTCTGGGCA AGCAACT  AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAATCTAGC AGAGTTT  TCTTCTGCCT CCATGTCTG CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGGTG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		AAGCTGAAAT	ATATGCATGT	AAAAACTTTG	ACATCTTTTT	TTTTAATTT	CCACTTTCTT	1320
CCAACCCCAA ATTIGTCTAA GCACTGGCCA GTCTGTTGTG GGCATTGTTT TCTACAA  AATTCTGGGT TTTTTTCTTC TTTCTTTAAA CATAGAGGTA CCACCACAAG GGATGCC  CTCTCTCGCA GCTCTTGAAA GCACTGTTT GAGGGAAAGG TCTCTGGGCA AGCAAGT  TATTTGGATT GCTTGCTTCC CTTTTTCCAC CTGGGACATT GYAATCATAA AATAACA  AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAATCTAGC AGAGTTT  CTTCTCGCCT CCATGCTGT CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA  GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA  GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA	10	CTTAACTTTA	CTTCTCTTTT	TGTCCCCCC	CCATCTTACA	GAAGTTGAGG	CCAAGGGAGA	1380
AATTCTGGGT TTTTTCTTC TTTCTTTAAA CATAGAGGTA CCACCACAAG GGATGCC CTCTCTCGCA GCTCTTGAAA GCATCTGTTT GAGGGAAAGG TCTCTGGGCA AGCAAGT  TATTTGATT GCTTGCTTCC CTTTTTCCAC CTGGGACATT GYAATCATAA AATAACA AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGT GAAATCTAGC AGAGTTI  TCTTCTGCCT CCATGTCTGT CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTACTATTC TTAGTTTGTA AATTCTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		ATGGTAGGCA	CAGAAGAAAC	ATGGCAAACT	GCTCTGTGCT	TTCAAACCAA	AGTGTTCCCC	1440
AATTCTGGGT TTTTTTCTTC TTTCTTTAAA CATAGAGGTA CCACCACAG GGATGCC CTCTCTCGCA GCTCTTGAAA GCATCTGTTT GAGGGAAAGG TCTCTGGGCA AGCAAGT  20 TATTTGGATT GCTTGCTTCC CTTTTTCCAC CTGGGACATT GYAATCATAA AATAACA AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAATCTAGC AGAGTTT  TCTTCTCCCT CCATGTCTGT CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  30 TGGACTTCTC ATCTAAAAGG TTAGTGGCTT TTGCTTGGGA TCAGTGCTCT CTATTGA TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGGATG GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAATT AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA	15	CCAACCCCAA	ATTTGTCTAA	GCACTGGCCA	GTCTGTTGTG	GGCATTGTTT	TCTACAACCA	1500
20 TATTTGGATT GCTTGCTTCC CTTTTTCCAC CTGGGACATT GYAATCATAA AATAACA AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAATCTAGC AGAGTTI  25 TCTTCTGCCT CCATGTCTGT CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  30 TGGACTTCTC ATCTAAAAGG TTAGTGGCTT TTGCTTGGGA TCAGTGCTCT CTATTGA TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		AATTCTGGGT	TTTTTTCTTC	TTTCTTTAAA	CATAGAGGTA	CCACCACAAG	GGATGCCCTA	1560
AATTCCAAAC CTCAAAAACT ATTATGGCCT GAGCACAGCT GAAACTCAGC AGAGTTI  TCTTCTGCCT CCATGTCTGT CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		CTCTCTCGCA	GCTCTTGAAA	GCATCTGTTT	GAGGGAAAGG	TCTCTGGGCA	AGCAAGTGGT	1620
25  TCTTCTGCCT CCATGTCTGT CACTTATAAT TCAGGTTCTG CTGTTGGCTT CAGAACA  GCAGAAGAAT CGTTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA  GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  30  TGGACTTCTC ATCTAAAAAG TTAGTGGCTT TTGCTTGGGA TCAGTGCTCT CTATTGA  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA	20	TATTTGGATT	GCTTGCTTCC	CTTTTTCCAC	CTGGGACATT	GYAATCATAA	AATAACAGTA	1680
GCAGAAGAAT COTTTATCC TAGTTATTCC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  TGACTTCTC ATCTAAAAGG TTAGTGGCTT TTGCTTGGGA TCAGTGCTCT CTATTGA  TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAAATA AAAACACTGT TGACAAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA		AATTCCAAAC	CTCAAAAACT	ATTATGGCCT	GAGCACAGCT	GAAATCTAGC	AGAGTTTAAC	1740
GCAGAAGAAT CGTTTATGC TAGTTATTGC ATTCATGGTT GAAACTCAAC TTAGGGA GGTTCCAATG TATTAAGCAA TGGGCTGCTT CTCCCCAATC CTCCCTAACA ATTCGTT  TGACTTCTC ATCTAAAAGG TTAGTGGCTT TTGCTTGGGA TCAGTGCTCT CTATTGA TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA	25	TCTTCTGCCT	CCATGTCTGT	CACTTATAAT	TCAGGTTCTG	CIGITGGCTT	CAGAACATGA	1800
TOTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		GCAGAAGAAT	CGTTTTATGC	TAGTTATTGC	ATTCATGGTT	GAAACTCAAC	TTAGGGAAAG	1860
TCTTGCTGGT CTCCAGACAC ATTCCTGTTG CATTAGACT TGAAAGACTT GTAGATG  GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		GGTTCCAATG	TATTAAGCAA	TEGECTECTT	CTCCCCAATC	CTCCCTAACA	ATTCGTTGTG	1920
GATGTTCAGG CACAGGATGC TGAAAGCTAT GTTACTATTC TTAGTTTGTA AATTGTC  TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA	30	TGGACTTCTC	ATCTAAAAGG	TTAGTGGCTT	TTGCTTGGGA	TCAGTGCTCT	CTATTGATGT	1980
TIGATACCAT CATCITGITT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA AAAAAA		TCTTGCTGGT	CTCCAGACAC	ATTCCTGTTG	CATTAAGACT	TGAAAGACTT	GTAGATGTGT	2040
TTGATACCAT CATCTTGTTT TCTTTTTGTA GGTATAAATA AAAACACTGT TGACAAT  AAAAAAAAAA AAAAAAAAA AAAAAAAAAA AAAAAA	35	GATGTTCAGG	CACAGGATGC	TGAAAGCTAT	GTTACTATTC	TTAGTTTGTA	AATTGTCCTT	2100
(2) INFORMATION FOR SEQ ID NO: 104:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1351 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT		TTGATACCAT	CATCTTGTTT	TCTTTTTGTA	GGTATAAATA	AAAACACTGT	TGACAATAAA	2160
(2) INFORMATION FOR SEQ ID NO: 104:  45  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1351 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  50  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT		AAAAAAAA	ААААААААА	АААААААА	ааааааааа	АААААААА	ААААААА	2218
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1351 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT	40		•					
(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1351 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT		(2) TNISODMA	TON FOR CE	O TD NO. 10				
(A) LENGTH: 1351 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT								
(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear  (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT	43	(i)						
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT			(B) TYPE	E: nucleic a	acid			
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:  CTTCACAGAC TGACAGAATG GTTTGTTTT GTTTTGTTTT			· ·	•				
CTTCACAGAC TGACAGAATG GTTTTGTTTT GTTTTGTTTT	50	(vi)	SECTIONS D	PCCBTDMTON.	CDO TD NO	104	•	
TOGACTCTAG CTCTGTCACC CAGGCTGGAG TGCAGTGGTG CGATCTCGGC TCACTGCT  CTCCGCCTCC CGGGTTCTCA CCATTCTCCT GCCTCAGCCT CCCGAGTAGC TGGGACTA  GGCGCCCACC ACCACGCCCG GCTAATTTTT TGTATTTTTT AGTAGAGACG GGGTTTCT			•					
CTCCGCCTCC CGGGTTCTCA CCATTCTCCT GCCTCAGCCT CCCGAGTAGC TGGGACTY  GGCGCCCACC ACCACGCCCG GCTAATTTTT TGTATTTTTT AGTAGAGACG GGGTTTC		CTTCACAGAC (	TGACAGAATG	GTTTTGTTTT	CTTTTCTTTT	GITTIGITTI	GTTTTTGAGA	60
GGCGCCCACC ACCACGCCCG GCTAATTTTT TGTATTTTTT AGTAGAGACG GGGTTTC	55	TGGACTCTAG (	CTCTGTCACC (	CAGGCTGGAG	TGCAGTGGTG	CGATCTCGGC	TCACTGCAAG	120
GGCGCCCACC ACCACGCCCG GCTAATTTTT TGTATTTTTT AGTAGAGACG GGGTTTCI		CTCCGCCTCC (	CGGGTTCTCA (	CCATTCTCCT	GCCTCAGCCT	CCCGAGTAGC	TGGGACTACA	180
• -	60	GGCGCCCACC 1	ACCACGCCCG (	GCTAATTTTT '	TGTATTTTTT .	AGTAGAGACG	GGGTTTCACC	240

	ATGTTAGCCA GGATGGTCTC GATCTCCTGA CCTCGTGATC CGCCCGCYTC GGCCTCCCAA	300
	AGTGCTGGGA TTACAGGCGT GAGCCACCGT GCCTGCCCCA GAATGGTTTT TAAAGCCACA	360
5	GTTGAGARGC CACCCATTGC CCGGCGCCTG GACAGTGATC ATCTTGTTCA TCTTGTTCAG	420
	TCCTTTCTTG TGTGATTGGA ATTATTCATC CCCTTTGAAA GATGAGAAGG TTGAGATGCA	480
10	AAGAGTCTAC CTTTCCAAGT TCTCACTGCT GGAAAGARCT AGAAGCACAG TTCAAAGTTC	540
	TGGNTTCTGG ACTCTGCAGT CCAGGTYTCC CTTYTCCCAC TTGCCTACCC TCAATGCCAC	600
	ACTGTTTTTG AAGTGGCCCA TAACTTGAAG GRAAAGTTTA AAGACAGTTC AATTTAATCA	660
15	TCAGRATGCA TTCTTTTTT TTTCGGARAC GGAKTTTCAC TCTTGCTGCC CASGCTGGAG	720
	TGCAATGGTG CAATGATCTC GGCTCACTGC AACCTATGCC TCCTGGGTTC AAGNGATTAT	780
20	CCAGCCTCAG CCTCCCGAGT AGCTGGGATT ATGGGCGCCC ACCACCATGC CCAGCTAATT	840
	TTTGTATTTT TTTTTTAGT AGAGATGGGG TTTCGCCAGG TTGGCCAGGC TGKTCTTGTG	900
	AAYTCCTGGC YTCAGGTGAT YTGCCCACYT CATCYTCCAA AAGTGCTGGG ATTACAGGCA	960
25	TGAGCCACTG CGCCTGGCYT CAGAATGCAT TCTTACACAT CTATCCTAGA CATTTATAAG	1020
	CACTCTAATG GATAACAATC CAAGAATAAA TGATTGTAAA AGATGATGCC GAAGAGTTGA	1080
30	TGTCAATCTT TTTTTCCTAA GAAAAAAAGT CCGCGAGTAT TAAATATTTA GATCAATGTT	1140
	TATAAAATGA TTACTTTGTA TATCTCATTA TTCCTATTTT GGAATAAAAA CTGACCTTCT	1200
	TTAATCATAT ACTTGTCTTT TGTAAATAGC AGCTTTTGTG TCATTCTCCC CACTTTATTA	1260
35	GTTAATTTAA ATTGGAAAAA ACCCTCAAAC TAATATTCTT GTCTGTTCCA GTCTTATAAA	1320
	TAAAACTTAT AATGCATGTA AAAAAAAAAA A	1351
40		
	(2) INFORMATION FOR SEQ ID NO: 105:	
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2066 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	<i>,</i>
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 105:	•
	GGCACGAGGC GGCGGAGGGC CACAATCACA GCTCCGGGCA TTGGGGGAAC CCGAGCCGGC	60
55	TGCGCCGGGG GAATCCGTGC GGGCGCCTTC CGTCCCGGTC CCATCCTCGC CGCGCTCCAG	120
<i>JJ</i>	CACCTCTGAA GTTTTGCAGC GCCCAGAAAG GAGGCGAGGA AGGAGGGAGT GTGTGAGAGG	180
	AGGGAGCAAA AAGCTCACCC TAAAACATTT ATTTCAAGGA GAAAAGAAAA	240
60	CAAAAATGGC TGGGGCAATT ATAGAAAACA TGAGCACCAA GAAGCTGTGC ATTGTTGGTG	300

	GGATTCTGCT	CGIGITCCAA	ATCATCGCCT	TTCTGGTGGG	AGGCTTGATT	GCTCCAGGGC	360
5	CCACAACGGC	AGTGTCCTAC	ATGTCGGTGA	AATGTGTGGA	TGCCCGTAAG	AACCATCACA	420
,	AGACAAAATG	GTTCGTGCCT	TGGGGACCCA	ATCATTGTGA	CAAGATCCGA	GACATTGAAG	480
	AGGCAATTCC	AAGGGAAATT	GAAGCCAATG	ACATCGTGTT	TTCTGTTCAC	ATTCCCCTCC	540
0	CCCACATGGA	GATGAGTCCT	TGGTTCCAAT	TCATGCTGTT	TATCCTGCAG	CTGGACATTG	600
٠	CCTTCAAGCT	AAACAACCAA	ATCAGAGAAA	ATGCAGAAGT	CTCCATGGAC	GTTTCCCTGG	660
5	CTTACCGTGA	TGACGCATTT	GCTGAGTGGA	CTGAAATGGC	CCATGAAAGA	GTACCACGGA	720
	AACTCAAATG	CACCTTCACA	TCTCCCAAGA	CTCCAGAGCA	TGAGGGCCGT	TACTATGAAT	780
	GTGATGTCCT	TCCTTTCATG	GAAATTGGGT	CTGTGGCCCA	TAAGTTTTAC	СТТТТАААСА	840
20	TCCGGCTGCC	TGTGAATGAG	AAGAAGAAAA	TCAATGTGGG	AATTGGGGAG	ATAAAGGATA	900
	TCCGGTTGGT	GGGGATCCAC	CAAAATGGAG	GCTTCACCAA	GGTGTGGTTT	GCCATGAAGA	960
25	CCTTCCTTAC	GCCCAGCATC	TTCATCATTA	TGGTGTGGTA	TTGGAGGAGG	ATCACCATGA	1020
	TGTCCCGACC	CCCAGTGCTT	CTGGAAAAAG	TCATCTTTGC	CCTTGGGATT	TCCATGACCT	1080
	TTATCAATAT	CCCAGTGGAA	TGGTTTTCCA	TCGGGTTTGA	CTGGACCTGG	ATGCTGCTGT	1140
30	TTGGTGACAT	CCGACAGGGC	ATCTTCTATG	CGATGCTTCT	GTCCTTCTGG	ATCATCTTCT	1200
	GTGGCGAGCA	CATGATGGAT	CAGCACGAGC	GGAACCACAT	TGCAGGGTAT	TGGAAGCAAG	1260
35	TCGGACCCAT	TGCCGTTGGC	TCCTTCTGCC	TCTTCATATT	TGACATGTGT	GAGAGAGGG	1320
	TACAACTCAC	GAATCCCTTC	TACAGTATCT	GGACTACAGA	CATTGGAACA	GAGCTGGCCA	1380
	TGGCCTTCAT	CATCGTGGCT	GGAATCTGCC	TCTGCCTCTA	CTTCCTGTTT	CTATGCTTCA	1440
10	TGGTATTTCA	GCTCTTTCGG	AACATCAGTG	GGAAGCAGTC	CAGCCTGCCA	GCTATGAGCA	1500
	AAGTCCGGCG	GCTACACTAT	GAGGGGCTAA	TTTTTAGGTT	CAAGTTCCTC	ATGCTTATCA	1560
15	CCTTGGCCTG	CGCTGCCATG	ACTGTCATCT	TCTTCATCGT	TAGTCAGGTA	ACGGAAGGCC	1620
	ATTGGAAATG	GGGCGCGTC	ACAGTCCAAG	TGAACAGTGC	CTTTTTCACA	GGCATCTATG	1680
	GGATGTGGAA	TCTGTATGTC	TTTGCTCTGA	TGTTCTTGTA	TGCACCATCC	CATAAAAACT	1740
50	ATGGAGAAGA	CCAGTCCAAT	GGAATGCAAC	TCCCATGTAA	ATCGAGGGAA	GATTGTGCTT	1800
	TGTTTGTTTC	GGAACTTTAT	CAAGAATTGT	TCAGCGCTTC	GAAATATTCC	TTCATCAATG	1860
55	ACAACGCAGC	TTCTGGTATT	TGAGTCAACA	AGGCAACACA	TGTTTATCAG	CTTTGCATTT	1920
-	GCAGTTGTCA	CAGTCACATT	GATTGTACTT	GTATACGCAC	ACAAATACAC	TCATTTAGCC	1980
	TTTATCTCAA	AATGTTAAAT	ATAAGGAAAA	AAGCGTCAAC	AATAAATATT	CTTGAGTATA	2040
60	АААААААА	ААААААААА	ААААА	`			2066

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)	(2)	INFORMATION	FOR	SEQ	ID	NO:	106:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1705 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(xi)	SEQUENCE I	DESCRIPTION	SEQ ID NO	: 106:		
15	AATTCGGCAK	AGGGCAGCTG	TCGGCTGGAA	GGAACTGGTC	TGCTCACACT	TGCTGGCTTG	60
	CGCATCAGGA	CTGGCTTTAT	CTCCTGACTC	ACGGTGCAAA	GGTGCACTCT	GCGAACGTTA	120
20	AGTCCGTCCC	CAGCGCTTGG	AATCCTACGG	CCCCCACAGC	CGGATCCCCT	CAGCCTTCCA	180
20	GGTCCTCAAC	TCCCGYGGAC	GCTGAACAAT	GGCCTCCATG	GGGCTACAGG	TAATGGGCAT	240
	CGCGCTGGCC	GTCCTGGGCT	GGCTGGCCGT	CATGCTGTGC	TGCGCGCTGC	CCATGTGGCG	300
25	CGTGACGGCC	TTCATCGGCA	GCAACATTGT	CACCTCGCAG	ACCATCTGGG	AGGGCCTATG	360
	GATGAACTGC	GTGGTGCAGA	GCACCGGCCA	GATGCAGTGC	AAGGTGTACG	ACTCGCTGCT	420
30	GGCACTGCCG	CAGGACCTGC	AGGCGGCCCG	CCCCTCCTC	ATCATCAGCA	TCATCGTGGC	480
	TGCTCTGGGC	GTGCTGCTGT	CCCTGGTGGG	GGGCAAGTGT	ACCAACTGCC	TGGAGGATGA	540
	AAGCGCCAAG	GCCAAGACCA	TGATCGTGGC	GGCGTGGTG	TTCCTGTTGG	CCGGCCTTAT	600
35	GGTGATAGTG	CCGGTGTCCT	GGACGGCCCA	CAACATCATC	CAAGACTTCT	ACAATCCGCT	660
	CCTCCCTCC	GGGCAGAAGC	GGGĄGATGGG	TGCCTCGCTC	TACGTCGGCT	GGGCCGCCTC	720
40	CGGNCTGCTG	CTCCTTGGCG	GGGGGCTGCT	TTGCTGCAAC	TGTCCACCCC	GCACAGACAA	780
	GCCTTACTCC	GCCAAGTATT	CTGCTGCCCG	CTCTGCTGCT	GCCAGCAACT	ACGTGTAAGG	840
	TGCCACGGCT	CCACTCTGTT	CCTCTCTGCT	TIGTICTICC	CTGGACTGAG	CTCAGCGCAG	900
45	GCTGTGACCC	CAGGAGGCCC	CTGCCACGGG	CCACTGGCTG	CTGGGGACTG	GGGACTGGGC	960
	AGAGACTGAG	CCAGGCAGGA	AGGCAGCAGC	CTTCAGCCTC	TCTGGCCCAC	TCGGACAACT	1020
50	TCCCAAGGCC	GCCTCCTGCT	AGCAAGAACA	GAGTCCACCC	TCCTCTGGAT	ATTGGGGAGG	- 1080
	GACGGAAGTG	ACAGGGTGTG	GTGGTGGAGT	GGGGAGCTGG	CTTCTGCTGG	CCAGGATGGC	1140
	TTAACCCTGA	CTTTGGGATC	TGCCTGCATC	GGTGTTGGCC	ACTGTCCCCA	TTTACATTTT	1200
55	CCCCACTCTG	TCTGCCTGCA	TCTCCTCTGT	TGCGGGTAGG	CCTTGATATC	ACCTCTGGGA	1260
	. CTGTGCCTTG	CTCACCGAAA	CCCGCGCCCA	GGAGTATGGC	TGAGGCCTTG	CCCACCCACC	1320
60	TGCCTGGGAA	GTGCAGAGTG	GATGGACGGG	TTTAGAGGGG	AGGGGCGAAG	GTGCTGTAAA	1380

WO 98/54963 PCT/US98/11422

358

	CAGGTTTGGG	CAGTGGTGGG	GGAGGGGGCC	AGAGAGGCGG	CTCAGGTTGC	CCAGCTCTGT	1440
	GGCCTCAGGA	CTCTCTGCCT	CACCCGCTTC	AGCCCAGGGC	CCCTGGAGAC	TGATCCCCTC	1500
5	TGAGTCCTCT	GCCCCTTCCA	AGGACACTAA	TGAGCCTGGG	AGGGTGGCAG	GGAGGAGGGG	1560
	ACAGCTTCAC	CCTTGGAAGT	CCTGGGGTTT	TTCCTCTTCC	TTCTTTGTGG	TTTCTGTTTT	1620
10	GTAATTTAAG	AAGAGCTATT	CATCACTGTA	ATTATTATTA	TTTTCTACAA	TAAATGGGAC	1680
10	CTGTGCACAG	GRAAAAAAAA	AAAAG				1705

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#### (2) INFORMATION FOR SEQ ID NO: 107:

### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1167 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 107:

TGCAGGAATT CGGCAGAGGT TTTCCGCTAG ACTCTGGCAG TTGGTGAGCA TCATGGCAAC 60 CGTTACAGCC ACAACCAAAG TCCCGGAGAT CCGTGATGTA ACAAGGATTG AGCGAATCGG 120 TGCCCACTCC CACATCCGGG GACTGGGGCT GGACGATGCC TTGGAGCCTC GGCAGGCTTC 180 GCAAGGCATG GTGGGTCAGC TGGCGGCACG GCGGCGGCT GGCGTGGTGC TGGAGATGAT 240 CCGGGAAGGG AAGATTGCCG GTCGGGCAGT CCTTATTGCT GGCCAGCCGG GCACGGGGAA 300 GACGGCCATC GCCATGGGCA TGGCGCAGGC CCTGGGCCCT GACACGCCAT TCACAGCCAT 360 CGCCGGCAGT GAAATCTTCT CCCTGGAGAT GAGCAAGACC GAGGCGCTGA CGCAGGCCTT 420 CCGCCGTCC ATCGCCGTTC GCATCAAGGA GGAGACGGAG ATCATCGAAG GGGAGGTGGT 480 GGAGATCCAG ATTGATCGAC CAGCAACAGG GACGGGCTCC AAGGTGGGCA AACTGACCCT 540 CAAGACCACA GAGATGGAGA CCATCTACGA CCTGGGCACC AAGATGATTG AKTCCCTGAC 600 CAAGGACAAG GTCCAGGCCG GGGACGTGAT CACCATCGAC AAGGCGACGG GCAAGATCTC 660 CAAGCTGGGC CGCTCCTTCA CACGCGCCCG CGAACTACGA CGCTATGGGC TCCCAGACCA 720 AGTTCGTGCA GTGCCCAGAT GGGGAGCTCC AGAAACGCAA GGAGGTGGTG CACACCGTGT 780 CCCTGCACGA GATCGACGTC ATCAACTCTC GCACCCAGGG CTTCCTGGCG CTCTTCTCAG 840 GTGACACAGG GGAGATCAAG TCAGAAGTCC GTGAGCAGAT CAATGCCAAG GTGGCTGAGT 900 GGCGCGAGGA GGGCAAGGCG GAGATCATCC CTGGAGTGCT GTTCATCGAC GAGGTCCACA 960 TGCTGGACAT CGAGAGCTTC TCCTTCCTCA ACCGGGCCCT GGAGAGTGAC ATGGCGCCTG 1020 TCCAGCAGGT CTATGGGGAT GCCGTGAGGG CTCTGGTAGC TGGTGCCCCG GATTCGCGTG 1080

WO 98/54963 PCT/US98/11422

	ATGCCACGGT TGGTGGCCTC GTGCCGAATT CCTGCAGCCC GGGGGATCCA CTAGTTCTAG	1140
5	AGCGGCCGCC ACCGCGGTGG ANCTCCN	1167
10	(2) INFORMATION FOR SEQ ID NO: 108:	•
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1907 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 108:	
20	GGCACAGGGG AATCATCGTG TGATGTGTGT GCTGCCTTTG TGAGTGTGTG GAGTCCTGCT	60
20	CAGGIGITAG GTACAGIGIG TITGATCGIG GIGGCTIGAG GGGAACCCTI GITCAGAGCT	120
	GTGACTGCGG CTGCACTCAG AGAAGCTGCC CTTGGCTGCT CGTAGCGCCG GGCCTTCTCT	180
25	CCTCGTCATC ATCCAGAGCA GCCAGTGTCC GGGAGGCAGA AGGTACCGGG GCAGCTACTG	240
	GAGGACTGTG CGGGCCTGCC TGGGCTGCCC CCTCCGCCGT GGGGCCCTGT TGCTGCTGTC	300
30	CATCTATTTC TACTACTCCC TCCCAAATGC GGTCGGCCCG CCCTTCACTT GGATGCTTGC	360
	CCTCCTGGGC CTCTCGCAGG CACTGAACAT CCTCCTGGGC CTCAAGGGCC TGGCCCCAGC	420
	TGAGATCTCT GCAGTGTGTG AAAAAGGGAA TTTCAACGTG GCCCATGGGC TGGCATGGTC	480
35	ATATTACATC GGATATCTGC GGCTGATCCT GCCAGAGCTC CAGGCCCGGA TTCGAACTTA	540
	CAATCAGCAT TACAACAACC TGCTACGGGG TGCAGTGAGC CAGCGGCTGT ATATTCTCCT	. 600
40	CCCATTGGAC TGTGGGGTGC CTGATAACCT GAGTATGGCT GACCCCAACA TTCGCTTCCT	660
	GGATAAACTG CCCCAGCAGA CCGGTGACCG TGCTGGCATC AAGGATCGGG TTTACAGCAA	720
	CAGCATCTAT GAGCTTCTGG AGAACGGGCA GCGGGGGGGC ACCTGTGTCC TGGAGTACGC	780
45	CACCCCCTTG CAGACTTTGT TTGCCATGTC ACAATACAGT CAAGCTGGCT TTAGCGGGGA	840
	GGATAGGCTT GAGCAGGCCA AACTCTTCTG CCGGACACTT GAGGACATCC TGGCAGATGC	900
50	CCCTGAGTCT CAGAACAACT GCCGCCTCAT TGCCTACCAG GAACCTGCAG ATGACAGCAG	960
	CTTCTCGCTG TCCCAGGAGG TTCTCCGGCA CCTGCGGCAG GAGGAAAAGG AAGAGGTTAC	1020
	TOTGGGCAGC TTGAAGACCT CAGCGGTGCC CAGTACCTCC ACGATGTCCC AAGAGCCTGA	
55	GCTCCTCATC AGTGGAATGG AAAAGCCCCT CCCTCTCCGC ACGGATTTCT CTTGAGACCC	
	AGGGTCACCA GGCCAGAGCC TCCAGTGGTC TCCAAGCCTC TGGACTGGGG GCTCTCTTCA	•
	CTCCCTCAAT CTCCACACA CCTATTTCCT TCCACACCCG CCCTTCCAGG GAAGGGTCCA	1260

WO 98/54963 PCT/US98/11422

360

•	GGACTTGACA	TCTTAAGATG	CGTCTTGTCC	CCTTGGGCCA	GTCATTTCCC	CTCTCTGAGC	. 1320
	CTCGGTGTCT	TCAACCTGTG	AAATGGGATC	ATAATCACTG	CCTTACCTCC	CTCACGGTTG	1380
5	TTGTGAGGAC	TGAGTGTGTG	GAAGTTTTTC	ATAAACTTTG	GATGCTAGTG	TACTTAGGGG	1440
	GTGTGCCAGG	TGTCTTTCAT	GGGGCCTTCC	AGACCCACTC	CCCACCCTTC	TCCCCTTCCT	1500
10	TTGCCCGGGG	ACGCCGAACT	CTCTCAATGG	TATCAACAGG	CTCCTTCGCC	CTCTGGCTCC	<sup>-</sup> 1560
10	TGGTCATGTT	CCATTATTGG	GGAGCCCCAG	CAGAAGAATG	GAGAGGAGGA	GGAGGCTGAG	1620
	TTTGGGGTAT	TGAATCCCCC	GGCTCCCACC	CTGCAGCATC	AAGGTTGCTA	TGGACTCTCC	1680
15	TGCCGGGCAA	CTCTTGCGTA	ATCATGACTA	TCTCTAGGAT	TCTGGCACCA	CTTCCTTCCC	1740
	TGGCCCCTTA	AGCCTAGCTG	TGTATCGGCA	CCCCACCCC	ACTAGAGTAC	TCCCTCTCAC	1800
20	TTGCGGTTTC	CTTATACTCC	ACCCCTTTCT	CAACGGTCCT	TTTTTAAAGC	ACATCTCAGA	1860
20	ТТАААААААА	АААААААА	АААААААА	AAAAAAAGGG	CGGCCGC		1907

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## (2) INFORMATION FOR SEQ ID NO: 109:

## (i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 611 base pairs (B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 109:

33	ATGAATTAAC	GCCAAGCTNT	NAATAGGGAC	TCACTATGGG	GGAAAGNTGG	GTAACGCCTG	60
	CAGGTACCGT	TCCGGAATTC	CCGGGTCGAC	CCACGCGTCC	GATGGGGCTT	TAGTAAATCA	120
40	GGCTTGCAGG	CTCAAAGCTG	CAATCTGCCC	ACTCTCAGGT	ACTGAGACTT	TGTGGGCCTC	180
45	AGACACCAGG	AAGAAAGTTG	GGATACAGTC	ATTTGAGTTA	AAAAGGGAAT	GACCCCTCAG	240
15	AAACCCGCAT	TAGCAGTGTT	ACTCTTGGAA	GTGCCTTTAC	TTTTAACGCT	CTCTGTTCTG	300
40	AAAAAGAGGT	GTTTGGTTAC	GTGTGAGCCA	ACATCACGTT	TTGTTAGCTG	TGATTTACCT	360
•	TIGICCGITT	AAAAGACTTC	ACGGAGCCAT	TCTGTATACA	AGGTGTGCTC	TITCCAATGT	120 180 240 300
50	AGAAGGGGTT	ATGGAAAAGG	GTGCGATCCT	TTGCTGTAAA	CTGGAGAGAC	CAGTCCCAAA	480
	CAGAGGGGAA	TTTTAAGCCC	TTCTCATCAC	CCAATTGGAT	GTTTTTGCTT	ATAGCAAATT	540
55	CCTGCAAAAT	АААТАААТАА	ATATTTGCAA	ААСТААААА	АААААААА	АААААААА	600
23	GGGGGGNCCN	С					611

## (2) INFORMATION FOR SEQ ID NO: 110:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2632 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 110:							
10	TCCCAGCTCT	CAGGACAAGG	GCCCTGGGCG	ATCTTTTAAA	AAAGCCGATT	GGGTGTCTTT	60	
	CTAAAANTAC	AACCAGTACT	.TCATCGTCAA	GTTTCTGGGA	AGGGAGTCCC	CTCCAGATTC	120	
15	TCATGGAGTG	ACAAATCTTG	ACTCTTGCTC	CTGGAATTTT	TCAGGCCCAA	ACTAGCGTTT	180	
	CTACAATGAT	TTATTTGGCA	AATTTGTCTT	GATTATGGGT	GGCTGATGAG	GAACGTGCTT	240	
20	TTGTTAGGAA	CCGAAACTGG	GCGGCGGTGA	GGGCGTGTAC	GCAATGAGTC	CGGAAGAGGG	300	
20	TGAAATGCTT	TCGGTAGGCA	CTCCACGGCT	GTGAAGATGG	CGGCGGCTGC	GTGGCTTCAG	360	
	GTGTTGCCTG	TCATTCTTCT	GCTTCTGGGA	GCTCACCCGT	CACCACTGTC	GTTTTTCAGT	420	
25	GCGGGACCGG	CAACCGTAGC	TGCTGCCGAC	CGGTCCAAAT	GGCACATTCC	GATACCGTCG	480	
	GGGAAAAATT	ATTITAGITT	TGGAAAGATC	CTCTTCAGAA	ATACCACTAT	CTTCCTGAAG	540	
30	TTTGATGGAG	AACCTTGTGA	CCTGTCTTTG	AATATAACCT	GGTATCTGAA	AAGCGCTGAT	600	
	TGTTACAATG	AAATCTATAA	CTTCAAGGCA	GAAGAAGTAG	AGTTGTATTT	GGAAAAACTT	660	
	AAGGAAAAA	GAGGCTTGTC	TGGGAAATAT	CAAACATCAT	CAAAATTGTT	CCAGAACTGC	720	
35	AGTGAACTCT	TTAAAACACA	GACCTTTTCT	GGAGATTTTA	TGCATCGACT	GCCTCTTTTA	786	
	GGAGAAAAAC	AGGAGGCTAA	GGAÇAATGGA	ACAAACCTTA	CCTTTATTGG	AGACAAAACC	840	
40	GCAATGCATG	AACCATTGCA	AACTTGGCAA	GATGCACCAT	ACATTTTTAT	TGTACATATT	900	
	GGCATTTCAT	CCTCAAAGGA	ATCATCAAAA	GAAAATTCAC	TGAGTAATCT	TTTTACCATG	960	
	ACTGTTGAAG	TGAAGGGTCC	CTATGAATAC	CTCACACTTG	AAGACTATCC	CTTGATGATT	1020	
45	TTTTTCATGG	TGATGTGTAT	TGTATATGTC	CTCTTTCCTC	TTCTGTGGCT	GGCATGGTCT	1080	
	GCCTGCTACT	GGAGAGATCT	CCTGAGAATT	CAGTTTTGGA	TIGGIGCIGI	CATCTTCCTG	1140	
50	GGAATGCTTG	AGAAAGCTGT	CTTCTATGCG	GAATTTCAGA	ATATCCGATA	CAAAGGARAA	1200	
	TCTGTCCAGG	GTGCTTTGAT	CCTTGCAGAR	CTGCTTTCAG	CAGTGAAACG	CTCACTGGCT	1260	
	CGAACCCTGG	TCATCATAGT	CAGTCTGGGA	TATGGCATCG	TCAAGCCACG	CCTGGAGTCA	1320	
55	CTCTTCATAA	GGTTGTAGTA	GCAGRAGCCC	TCTATCTTTT	GTTCTCTGGC	ATGGAAGGGG	1380	
	TCCTCAGAGT	TACTGGGGCC	CAGACTGATC	TTGCTTCCTT	GCCTTTATC	CCCTTGGCTT	1440	
60	TCCTAGACAC	TGCCTTGTGC	TGGTGGATAT	TTATTAGCCT	GACTCAAACA	ATGAAGCTAT	1500	

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	TAAAACTTCG GAGGAACATT GTAAAACTCT CTTTGTATCG GCATTTCACC AACACGCTTA	1560
	TTTTGGCAGT GGCAGCATCC ATTGTGTTTA TCATCTGGAC AACCATGAAG TTCAGAATAG	1620
5	TGACATGTCA GTCGGACTGG CGGGAGCTGT GGGTAGACGA TGCCATCTGG CGCTTGCTGT	1680
	TCTCCATGAT CCTCTTTGTC ATCATGGTTC TCTGGCGACC ATCTGCAAAC AACCAGAGGT	1740
10	TTGCCTTTTC ACCATTGTCT GAGGAAGAGG AGGAGGATGA ACAAAAGGAG CCTATGCTGA	1800
	AAGAAAGCTT TGAAGGAATG AAAATGAGAA GTACCAAACA AGAACCCAAT GGAAATAGTA	1860
	AAGTTAACAA AGCACAGGAA GATGATTTGA AGTGGGTAGA AGAGAATGTT CCTTCTTCTG	1920
15	TGACAGATGT AGCACTTCCA GCCCTTCTGG ATTCAGATGA GGAACGAATG ATCACACACT	1980
	TTGAAAGGTC CAAAATGGAG TAAGGAATGG GAAGATTTGC AGTTAAAGAT GGCTACCATC	2040
20	AGGGAAGAGA TCAGCATCTG TGTCAGTCTT CTGTACGGCT CCATGGGATT AAAGGAAGCA	2100
	ATGACATCCT GATCTGTTCC TTGATCTTTG GGCATTGGAG TTGGCGAGAG GTGTCAGAAC	2160
	AAAGAGAACA TCTTACTGAA AACAAGTTCA TAAGATGAGA AAAATCTACG AGCTTCTTAT	2220
25	TTACAACACT GCTGCCCCCT TTCCTCCCAG ACTCTGACAT GGATGTTCAT GCAACTTAAG	2280
	TGTGTTGTTC CTGAACTTTC TGTAATGTTT CATTTTTTAA ATCTGACAAA CTAAAAAGTT	2340
30	TAACGTCTTC TAAAAGATTG TCATCAACAC CATAATATGT AATCTCCAGG AGCAACTGCC	2400
	TGTAATTTTT ATTTATTTAG GGAGTTACAT AGGTGATGGG GGAAATTGTT AACTACCTTT	2460
	CATTITCCTG GGAAGTCAAG GTTACATCTT GCAGAGGTTG TTTTGAGAAA AAAGGGCCCT	2520
35	TCTGAGTTAA GGAGCCATAG TTCTATCAAT GATCAAAAGA AAAAAAAAA AACTCGATCG	2580
	GCACGAGGGG GGGCCCGGTA CCCAATTCGC CCTATGGGAN TCGAATGAGA CC	2632
40		
	(2) INFORMATION FOR SEQ ID NO: 111:	
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2249 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	·
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 111:	
	GAATTCGGCA CGAGCTCACC GTGCTGCGTG ACACAAGGCC AGCCTGCGCC TACGAGCCCA	. 60
	TGGACTTIKT RATGGCCCTC ATCTACGACA TGGTACTGSW TGTGGTCACC CTGGGGCTGG	120
55	CCCTCTTCAC TCTGTGCGGC AAGTTCAAGA GGTGGAAGCT GAACGGGCC TTCCTCCTCA	180
	TCACAGCCTT CCTCTCTGTG CTCATCTGGG TGGCCTGGAT GACCATGTAC CTCTTCGGCA	240

ATGTCAAGCT GCAGCAGGGG GATGCCTGGA ACGACCCCAC CTTGGCCATC ACGCTGGCGG

	CCAGCGCTGG	GTCTTCGTCA	TCTTCCACGC	CATCCCTGAG	ATCCACTGCA	CCCTTCTGCC	360
5	AGCCCTGCAG	GAGAACACGC	CCAACTACTT	CGACACGTCG	CAGCCCAGGA	TGCGGGAGAC	420
3	GGCCTTCGAG	GAGGACGTGC	AGCTGCCGCG	GGCCTATATG	GAGAACAAGG	CCTTCTCCAT	480
	GGATGAACAC	AATGCAGCTC	TCCGAACAGC	AGGATTTCCC	AACGGCAGCT	TGGGAAAAAG	. 540
10	ACCCAGTGGC	AGCTTGGGGA	AAAGACCCAG	CGCTCCGTTT	AGAAGCAACG	TGTATCAGCC	600
	AACTGAGATG	GCCGTCGTGC	TCAACGGTGG	GACCATCCCA	ACTGCTCCGC	CAAGTCACAC	660
15	AGGAAGAMAC	CTTTGGTGAA	AGACTTTAAG	TTCCAGAGAA	TCAGAATTTC	TCTTACCGAT	720
	TTGCCTCCCT	GCTGTGTCT	TTCTTGAGGG	AGAAATCGGT	AACAGTTGCC	GAACCAGGCC	780
	GCCTCACAGC	CAGGAAATTT	GGAAATCCTA	GCCAAGGGGA	TTTCGTGTAA	ATGTGAACAC	840
20	TGACGAACTG	AAAAGCTAAC	ACCGACTGCC	CCCCCTCCC	CTGCCACACA	CACAGACACG	900
	TAATACCAGA	CCAACCTCAA	TCCCCGCAAA	CTAAAGCAAA	GCTAATTGCA	AATAGTATTA	960
25	GGCTCACTGG	AAAATGTGGC	TGGGAAGACT	GTTTCATCCT	CTGGGGGTAG	AACAGAACCA	1020
	AATTCACAGC	TGGTGGCCA	GACTGGTGTT	GGTTGGAGGT	GGGGGGCTCC	CACTCTTATC	1080
	ACCTCTCCCC	AGCAAGTGCT	GGACCCCAGG	TAGCCTCTTG	GAGATGACCG	TTGCGTTGAG	1140
30	GACAAATGGG	GACTTTGCCA	CCGGCTTTGC	CIGGIGGITT	GCACATTTCA	GGGGGGTCAG	1200
	GAGAGTTAAG	GAGGTTGTGG	GTGGGATTCC	AAGGTGAGGC	CCAACTGAAT	CGTGGGGTGA	1260
35	GCTTTATAGC	CAGTAGAGGT	GGAGGGACCC	TGGCATGTGC	CAAAGAAGAG	GCCCTCTGGG	1320
	TGATGAAGTG	ACCATCACAT	TTGGAAAGTG	ATCAACCACT	GTTCCTTCTA	TGGGGCTCTT	1380
	GCTCTAGTGT	CTATGGTGAG	AACACAGGCC	CCGCCCCTTC	CCTTGTAGAG	CCATAGAAAT	1440
40	ATTCTGGCTT	GGGCAGCAG	TCCCTTCTTC	CCTTGATCAT	CTCGCCCTGT	TCCTACACTT	1500
	ACGGGTGTAT	CTCCAAATCC	TCTCCCAATT	TTATTCCCTT	ATTCATTTCA	AGAGCTCCAA	1560
45	TGGGGTCTCC	AGCTGAAANS	CCCTCCGGGA	GGCAGGTTGG	AAGGCAGGCA	CCACGGCAGG	1620
	TTTTCCGCGA	TGATGTCACC	TAGCAGGGCT	TCAGGGGTTC	CCACTAGGAT	GCAGAGATGA	1680
-	CCTCTCGCTG	CCTCACAAGC	AGTGACACCT	CGGGTCCTTT	CCGTTGCTAT	GGTGAAAATT	1740
50	CCTGGATGGA	ATGGATCACA	TGAGGGTTTC	TTGTTGCTTT	TGGAGGGTGT	GGGGGATATT	1800
	TIGTITIGGT	TTTTCTGCAG	GTTCCATGAA	AACAGCCCTT	TTCCAAGCCC	ATTGTTTCTG	1860
55	TCATGGTTTC	CATCTGTCCT	GAGCAAGTCA	TICCITIGIT	ATTTAGCATT	TCGAACATCT	1920
<i></i>	CGCCATTCA	AAGCCCCCAT	GTTCTCTGCA	CTGTTTGGCC	AGCATAACCT	CTAGCATCGA	1980
	TTCAAAGCAG	AGTTTTAACC	TGACGGCATG	GAATGTATAA	ATGAGGGTGG	GTCCTTCTGC	2040
60	AGATACTCTA	ATCACTACAT	TGCTTTTTCT	ATAAAACTAC	CCATAAGCCT	TTAACCTTTA	2100

WO 98/54963 PCT/US98/11422

	AAGAAAAATG AAAAAGGTTA GTGTTTGGGG GCCGGGGGAG GACTGACCGC TTCATAAGCC	2160
5	AGTACGTCTG AGCTGAGTAT GTTTCAAFAA ACCTTTTGAT ATTTCTCAAA AAAAAAAAAA	2220
3	AAAAANCCCG GGGGGGGGCC CGGACCTGG	2249
10	(2)	
	(2) INFORMATION FOR SEQ ID NO: 112:	
	(i) SEQUENCE CHAPACTERISTICS:  (A) LENGTH: 2193 base pairs	
15	(3) TYPE: nucleic acid	
	(C) STRAIDENESS: double (D) TOPCLOGY: linear	
	(b) 15F01001. 121.ea.	
20	(xi) SEQUENCE DESCFIPTION: SEQ ID NO: 112:	
	GATACTATAA GGCAAGTGAC TCACGGGTGC GCCGTTAGAC TAGTGGATCC CGGGTGCAGG	60
	AATTOGGCAG AGCGCCGCCG GAGCCGAAGT GCTGGCGCCCC CCGCGGCCGC TGCCTCCGCG	120
25	GANCCCAAAA TCATGAAAST CACCGTGAAG ACCCCGAAGA AAAGGAGGAA TTCGCCGTGC	180
	CCGAGAATAG CTCCGTCCAG CAGTTTAAGG AAGAAATCTC TAAACGTTTT AAATCACATA	240
30	CTGACCAACT TGTGTTGATA TTTGCTGGAA AAATTTTGAA AGATCAAGAT ACCTTGAGTC	300
50	AGCATGGAAT TCATGATGGA CTTACTGTTC ACCTTGTCAT TAAAACACAA AACAGGCCTC	360
	AGGATCATTC AGCTCAGCAA ACAAATACAG CTGGAAGCAA TGTTACTACA TCATCAACTC	420
35	CTAATAGTAA CECTACATOT GGETCTGCTA CTAGCAACCC TTTTGGTTTA GGTGGCCTTG	480
	GGGGACTTGC AGGTCTGAST AGCTTGGGTT TGAATACTAC CAACTTCTCT GAACTACAGA	540
40	GTCAGATGCA GCGACAACIT TTGTCTAACC CTGAAATGAT GGTCCAGATC ATGGAAAAWC	600
10	CCYTTGTTCA GAGCATGCTC MTCAAATCCT GACCTGATGN AGACAGTTAA TTATGGCCAA	660
	TCCACAAATG CAGCAGTTGA TACAGAGAAA TCCCAGAAAT TAGTCATATG TTGAATAATC	720
45	CAGATATAAT GAGACAAACG TTGGAACTTG CCCAGGAATC CAGCAATGAT GCAGGAGATG	780
	ATGAGGAACC AGGACCGAGC TTTGAGCAAC CTAGAAAGCA TCCCAGGGGG ATATAATGCT	840
50	TTAAGGCGCA TGTACACAGA TATTCAGGAA CCAATGCTGA GTGCTGCACA AGAGCAGTTT	900
50	GGTGGTAATC CATTTGCTTC CTTGGTGAGC AATACATCCT CTGGTGAAGG TAGTCAACCT	960
	TCCCGTACAG AAAATAGAGA TCCACTACCC AATCCATGGG CTCCACAGAC TTCCCAGAGT	1020
55	TCATCAGCTT CCAGCGGCAC TGCCAGCACT GTGGGTGGCA CTACTGGTAG TACTGCCAGT	1080
	GGCACTTCTG GGCAGAGTAC TACTGCGCCA AATTTGGTGC CTGGAGTAGG AGCTAGTATG	1140
	TTCAACACAC CAGGAATSCA GAGSTTGTTG CAACAAATAA CTGAAAAACCC ACAACTTATG	1200

WO 98/54963 PCT/US98/11422

-	CAAAACATGT TGTCTGCCCC CTACATGAGA AGCATGATGC AGTCACTAAG CCAGAATCCT	1260
	GACCTTGCTG CACAGATGAT GCTGAATAAT CCCCTATTTG CTGGAAATCC TCAGCTTCAA	1320
5	GAACAAATGA GACAACAGCT CCCAACTTTC CTCCAACAAA TGCAGAATCC TGATACACTA	1380
-	TCAGCAATGT CAAACCCTAG AGCAATGCAG GCCTTGTTAC AGATTCAGCA GGGTTTACAG	1440
	ACATTAGCAA CGGAAGCCCC GGGCCTCATC CCAGGGTTTA CTCCTGGCTT GGGGGCATTA	1500
10	GGAAGCACTG GAGGCTCTTC GGGAACTAAT GGATCTAACG CCACACCTAG TGAAAACACA	1560
	AGTCCCACAG CAGGAACCAC TGAACCTGGA CATCAGCAGT TTATTCAGCA GATGCTGCAG	1620
15	CCTCTTCCTG GAGTAAATCC TCAGCTACAG AATCCAGAAG TCAGATTTCA GCAACAACTG	1680
	GAACAACTCA GTGCAATGGG ATTTTTGAAC CGTGAAGCAA ACTTGCAAGC TCTAATAGCA	1740
	ACAGGAGGTG ATATCAATGC AGCTATTGAA AGGTTACTGG GCTCCCAGCC ATCATAGCAG	1800
20	CATTTCTGTA TCTKGAAAAA ATGTAATTTA TTTTTGATAA CGGCTCTTAA ACTTTAAAAT	1860
	ACCIGCTITA TITICATTITG ACTOTICGAA TICTGTGCTG TIATAAACAA ACCCAATATG	1920
25	ATGCATTTTA AGGTGGAGTA CAGTAAGATG TGTGGGTTTT TCTGTATTTT TCTTTTCTGG	1980
	AACAGTGGGA ATTAAGGCTA CTGCATGCAT CACTTCTGCA TTTATTGTAA TTTTTTAAAA	2040
	ACATCACCTT TTATAGTTGG GTGACCAGAT TTTGTCCTGC ATCTGTCCAG TTTATTTGCT	2100
30	TTTTAAACAT TAGCCTATGG TAGTAATTTA TGTAGAATAA AAGCATTAAA AAGAAGCAAA	2160
	AAAAAAAAA AAAAATTCCT GCGCCCGCGA ATTCTTCT	2198
35	ARRAMANA ARRATICCI GEGEGEGEA ATTETTET	2170
55	•	
	(2) INFORMATION FOR SEQ ID NO: 113:	
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1043 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
45	(D) TOPOLOGY: linear	
73	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 113:	
	CTGAAGTGTA TGTGGTGAGG AAGAAGAGGC TCCTACTGTA GACAGCCTTG TTCTACAGAT	. 60
50	CCTCCCAGAA ATCTCTGGGC CAGGTGGAAC CCAGGGTCAG AGAGGGATGG GAGAGAGGTT	120
	TAATTITCCA TGATAAATAA AAATCTATAA AATAATAAAC AAGAGAAAAG AGATTGGAAA	180
55	CAGCCAGGIT GGAGCAGIGA GIGAGIAAGG AAACCIGGCI GCCCTCTCCA GATTCCCCAG	240
<i>J J</i>	GCTCTCAGAG AAGATCAGCA GAAAGTCTGC AAGACCCTAA GAACCATCAG CCCTCAGCTG	300
	CACCTCCTCC CCTCCAAGGA TGACAAAGGC GCTACTCATC TATTTGGTCA GCAGCTTTCT	360
60	TGCCCTAAAT CAGGCCAGCC TCATCAGTCG CTGTGACTTG GCCCAGGTGC TGCAGCTGGA	420

	RGACTTGGAT GGGTTTGAGG GTTACTCCCT GAGTGACTGG CTGTGCCTGG CTTTTGTGGA	480
5	AAGCAAGTTC AACATATCAA AGATWAATGA AAATGCAGAT GGAAGCTTTG ACTATGGSCT	540
5	CTTCCAGATC AACAGCCACT ACTGGTGCAA CRATTATAAG AGTTACTCGG AAAACCTTTG	600
	CCACGTAGAC TGTCAAGATC TGCTGAATCC CAACCTTCTT GCAGGCATCC ACTGCGCAAA	. 660
10	AAGGATTGTG TCCGGAGCAC GGGGGATGAA CAACTGGGTT AGAATGGAAG KTTGCACTGT	720
	TCAGGCCGGC CACTCTTCTA CTGGCTGACA GGATGCCGCC TGAGATKAAA CARGGTGCGG	780
15	GTGCACCGTG GARTCATTCC AAGACTCCTG TCCTCACTCA RGGATTCTTC ATTTCTTCTT	840
15	CCTACTGCCT CCACTTCATG TTATTTTCTT CCCTTCCCAT TTACAACTAA AACTGACCAG	900
	AGCCCCAGGA ATAAATGGTT TTCTTGGCTT CCTCCTTACT CCCATCTGGA CCCAGTCCCC	960
20	TEGTTCCTGT CTGTTATTTG TAAACTGAGG ACCACAATAA AGAAATCTTT ATATTTATCG	1020
	AAAAAAAAA AAAAAAAACT CGA	1043
25		
·	(2) INFORMATION FOR SEQ ID NO: 114:	·
30	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 703 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li><li>(D) TOPOLOGY: linear</li></ul>	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 114:	
	GAATTCGCCA CGAGTGCGCG GGCACCACGG CGGTTTTTCG ACGCTGGCGG TGGACGCACG	60
40	CASCATGGAC CACGGTTGCT GGGCGGATGG GGAGCGTCTA TGGTCAGTTG CCTTAGAAGT	120
70	GGTGAGATGG GAAGCTGCAG TTGGAAGACC CTGGAGGATG CCTGACAAGG GGATGTCTGA	180
	CACATGATTG GAGCTCTTTT TGAAATGTTT CTTGCCCTTC CTGGAGCAGA GGAGCCATTA	240
45	TITATGCAGG TACATCGAAG TCTTTTGACC TCCATACAGT GATTATGCTT GTCATCGCTG	300
	GTGGTATCCT GGCGGCCTTG CTCCTGCTGA TAGTTGTCGT GCTCTGTCTT TACTTCAAAA	360
50	TACACAACGC GCTAAAAGCT GCAAAGGAAC CTGAAGCTGT GGCTGTAAAA AATCACAACC	420
50	CAGACAAGGT GTGGTGGGCC AAGAACAGCC AGGCCAAAAC CATTGCCACG GAGTCTTGTC	480
	CTGCCCTGCA GTGCTGTGAA GGATATAGAA TGTGTGCCAG TTTTGATTCC CTGCCACCTT	540
55	GCTGTTGCGA CATAAATGAG GGCCTCTGAG TTAGGAAAGG TGGGCACAAA AATCTTCATG	600
	AGCAATACTT CTTAGTAGAT TGTTTTGTTA TTCAAATCAA GTTCTAGTGT TTTTATGTGA	660
	GATTATATAA TITACAGTGT TGTTTTATAT ACTTTTGAAT AAA	703

1440

5	(2) INFORMATION FOR SEQ ID NO: 115:	
J	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 3684 base pairs  (B) TYPE: nucleic acid	
10	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 115:	
15	GGCAGAGGGG GCATGAGCAG GAGGAGGATT ACCGCTACGA GGTGCTCACG GCCGAGCAGA	. 60
15	TTCTACAACA CATGGTGGNA ATGTATCCGG GAGGTCAACG AGGTCATCCA GAATCCAGCA	12
	ACTATCACAA GAATACTCCT TAGCCACTTC AATTGGGATA AAGAGAAGCT AATGGAAAGG	18
20	TACTITIGATG GAAACCIGGA GAAGCTCTIT GCTGAGTGTC ATGTAATTAA TCCAAGTAAA	. 24
	AAGTCTCGAA CACGCCAGAT GAATACAAGG TCATCAGCAC AGGATATGCC TTGTCAGATC	·30
25	TGCTACTTGA ACTACCCTAA CTCGTATTTC ACTGGCCTTG AATGTGGACA TAAGTTTTGT	36
23	ATGCAGTGCT GGAGTGAATA TTTAACTACC AAAATAATGG AAGAAGGCAT GGGTCAGACT	42
	ATTTCGTGTC CTGCTCATGG TTGTGATATC TTAGTGGATG ACAACACAGT TATGCGCCTG	48
30	ATCACAGATT CAAAAGTTAA ATTAAAGTAT CAGCATTTAA TAACAAATAG CTTTGTAGAG	54
	TGCAATCGAC TGTTAAAGTG GTGTCCTGCC CCAGATTGCC ACCATGTTGT TAAAGTCCAA	60
35	TATCCTGATG CTAAACCTGT TCGCTGCAAA TGTGGGCGCC AATTTTGCTT TAACTGTGGA	66
رې	GAAAATTGGC ATGATCCTGT TAAATGTAAG TGGTTAAAGA AATGGATTAA AAAGTGTGAT	72
	GATGACAGTG AAACCTCCAA TTGGATTGCA GCCAACACAA AGGAATGTCC CAAATGCCAT	78
40	GTCACAATTG AGAAGGATGG TGGTTGTAAT CACATGGTCT GTCGTAACCA GAATTGTAAA	84
	GCAGAGTTTT GCTGGGTGTG TCTTGGCCCA TGGGAACCAC ATGGATCTGC CTGGTACAAC	90
15	TGTAACCGCT ATAATGAGGA TGATGCAAAG GCAGCAAGAG ATGCACAGGA GCGATCTAGG	96
45	GCAGCCCTGC AGAGGTACCT GTTCTACTGT AATCGCTATA TGAACCACAT GCAGAGCCTG	102
	CGCTTTGAGC ACAAACTATA TGCTCAGGTG AAACAGAAAA TGGAGGAGAT GCAGCAGCAC	108
50	AACATGTCCT GGATTGAGGT GCAGTTCCTG AAGAAGGCAG TTGATGTCCT CTGCCAGTGT	114
	CGTGCCACAC TCATGTACAC TTATGTCTTC GCTTTCTACC TCAAAAAGAA TAACCAGTCC	120
ـــر ســر	ATTATCTTTG AGAATAACCA AGCAGATCTA GAGAATGCCA CAGAGGTGCT CTCGGGCTAC	126
55	CTTGAACGAG ATATTTCCCA AGATTCTCTG CAGGATATAA AGCAGAAAGT ACAAGACAAG	132
	TACACATACT CTCACACTCC ACCAACCCTT TIVETTACACC ATCTCATCA ACCCTATCAA	170

AAAGATCTGT GGGAGTACAT TGAGGACTGA GAATGGCCCT GCATAAAATG AACTCTGAAA

	ACTITACCAT	CTAGAGTGCT	' CATGCAATTA	AAACAAAACA	AACACAAACA	AGGAGGCACT	150
5	AAGCCTATTC	TGACACCACT	GGTCTGTAGT	ACCAGAATTG	TTTTGTTAAT	GGAAAGTTTA	156
•	AGTAAATTAT	ATTGTAATAA	AAAGGTAGAT	AAACCATTGT	' ACAACAGTAT	TCTAGGCCGC	162
	CAACAAAAGT	GTGACAGACA	CACTAAAAGC	CCTCCAACTT	TAACTTGTAA	CGTAGCTTCA	. 168
10	TTCTCAAAGC	TGACTCCTTT	TTTTTCTTT	TCCTTTTCCT	GAGTGTAGTA	CAGTTAAAAT	174
	TTCAAACAGC	TCCTTGACAC	TGCTTTTCAT	GTTCAAACCA	GCCATTTTGT	TGTACTTTGG	180
15	TAAAGGACCT	CTTCCCCTTC	CTCCCCTACA	CATACAGATA	CACCCACACA	CAGACTGACT	186
	CTCTTTCTCT	CATACCCCAA	GGTCATGAGT	GAATGATGCT	TAGTTCCTTG	TAAAGAAAAT	192
	CTTGGGATGG	GGAAAGGGGT	AGGCAGCAAG	AGGATTCAAC	AAACGAAAAA	САТАААААСТ	198
20	TTGTATATGA	CTTTTAAAAC	AAGAGGACAA	CACAGTATTT	TTCAAAATTG	TATATAGCGC	204
	ATATGCATGG	ACAAAGCAAG	CGTGGCACGT	GTTTGCATAA	TGTTTAATTA	СААААААТА	210
25	TTTATTCTTT	AAAAATCTTC	AAGATTATGT	CTATTTGCTG	TGCATTTTCT	TTCAGTTTGC	216
	TTATCTTTCC	CGGGTTGGGG	TTGGGATAAA	GGTGTGTCGG	TTTAGCACCT	CTGGAAGACC	222
	TATCTAGAGC	TCTTTCACTT	TCCTGAGGTT	ATTTTGCCCY	TTCTGGTGTT	GCTATCTCTG	2280
30	TTGCCGGCCA	TGGGCTNCAY	GCCTTGAATT	CCTGCTCTTG	ATCAGGGACA	AGGGAGGTCA	2340
	AGCTCTGACT	AATGCCATGA	CCTGATTAAG	GGGTACAGCA	GGGAGTTTTG	TTGCTACAGC	2400
35	TCATGAATTA	ACCTGTCCCA	ACCTAATCCC	CCTCCATGGC	ATCATGCCTC	TACCCAAGCC	2460
	TTTGTGTGCC	CATGTTATGC	ACACAGCTGT	AGGCATTCTT	AAGTCCCCTG	TCGCATCCAG	2520
	TGGAAGCATT	TTAAAATTTC	TTTTACTTTT	TGGTTTTCCC	TTAATTGCTG	CTTTTCAGAT	2580
40	TTTAGTTATG	GCTCGTCTGC	TCACCCCTTC	TCTACATTAG	GGTGTCAAAG	AGAATGTTTT	2640
	GCTTTAAATA	TAAATAGCCA	TTCATTTAGT	CTCAGATTGT	GAATTTAAAA	TGGTGGATAC	2700
45	CGAAATTGCT	TGTGTGTGTT	GCTGTGGGTT	TGGTTTGAAG	GCAAACACCC	CTAGAACATG	2760
	ATATTCCCAT	CTAGTGCATT	TAAATAGAAA	TCACTGAGTT	TGCTGCTTTT	TTATTGTCAG	2820
	CAGATAGGAG	AATTAATAAT	GCATTTTAGC	TGTGATGTCC	ATTTTTATGA	AATTCCTACT	2880
50	AAGAGCTATG	TTAAAAGTAA	AGGATGGTGG	TGGTTGTATT	AACTATATAC	CTGTTTAGGC	2940
	CATTCTGGCT	GTGGTATT <b>T</b>	TCAATAGGTC	AGCATCTGTA	AATCTGTCAG	TTTTATACAG	3000
55	GAGTGCAGAG	TGAACTAGGC	AACTAGATTA	AGAGGTCTAA	ATATGAAATA	CCAGTTGAGG	3060
	CTGAGGACCT	CTTCGTCTTC	CTTTAAATGT	CTTTTGCCTA	GGGAGTGTTT	ACCATTTGTG	3120
	AGGCAGCTTT	GTCTGCTCTT	ACACTGTACA	TCCTATTACT	CCATTGGGAA	GTAGGTTCAC	3180
60	TTTCCTCTCG	CCTTTTGCCT	AAGTTAGGCT	TTGCTGAATC	AACCCTACTT	TTCCTTTTAG	3240

WO 98/54963 PCT/US98/11422

	AAAAGGTTGT TACAGGAGAT TTACTGGCAA CTGTTCTTTT CCCATCAAAA ATCAGTGAAT	3300
5	GTTTGCTGAG TATAAATGCT GCTTCCTTAA ACCACTTGTC GCTTTAGGAT CAACTTTACC	3360
5	TGTACCTTTT CTCCTTTCCT CCCTTGCCAC CTCAGGTGCA AATCTGAACT CAGTGTCTGC	3420
	TTCTTCCATT TTCTCGTCTC TCTCCCCTCT TCCCCCATTA TCCATATGAC ATTATTTTAC	3480
10	TTCAAATGAC AGCATCAATC TTAAAAAGAT ATACATTAAA ACTAAGGAGT TTTTTTAAAG	3540
	AAAGCCTGAA TAAGTTCCTT TCCCTGGTAA CTTTGAAAAG CAGTCAGAGT TGCTATATAG	3600
15	ATATATGTGG CTCCTTTAAA ATGCTTTGTG TATGTGTGT GTTTAAAAAA AAAAAAAAA	3660
13	TTCGGGGGG GGCCCGGTNC CCAT	3684
20	(2) THEODYSTON FOR CTO TO NO. 116	
	(2) INFORMATION FOR SEQ ID NO: 116:	
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1965 base pairs	
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
30	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 116:	
	AAGAAAGGGT ATTAAAATTC TAGATCACAT ATGGACCCGG GAAGGTTTTT NACCCTCTGT	60
	TAGTGACATC GAGTCTCCCA CTAGACAAAA TAGGTGGAAA AATCTCTCGA GGGCTCACAT	120
35	TGTTTTGTCA TCTTCAGGAA AAACACCACC AGGCCATACC ACAGCCTGCC CAGTGAGGCG	180
	GTCTTTGCCA ACAGCACCGG GATGCTGGTG GTGGCCTTTG GGCTGCTGGT GCTCTACATC	240
40	CTTCTGGCTT CATCTTGGAA GCGCCCAGAG CCGGGGGATCC TGACCGACAG ACAGCCCCTG	300
	CTGCATGATG GGGAGTGAAG CAGCAGGAAG GGGCTCCCAA GAGCTCCTGG TGGTGCAGCC	360
	TOTGCTCCCC TCAGAAGCTC TGCTCTTCCC AGGGCTCCCG GCTGGTTTCA GCAGGCGACT	420
45	TICTTCCAAT GCTGGGCCCA GACTTCTTGC CTGGGTGCTG GCCTGCCCTC TCCGGNCCGC	480
	TIGCTGCCTG TCTGCTTTCC TIGGTGGYTT TGCTGGGTGC TGGGCCTGCC CTCTCCGGCC	540
50	GCTTGCTGCC TGTCTGCTTT CCTTGGTGGC TTTGCTGGGT GCTGGGCCTG CCTTCTCTGG	600
	CTGCTTGCTG CCTGTCTGCT TTCCTTGGTG GCTTTGGCTT CTGCACTCCT TGGCGTCASC	660
	TCTCAGGTCC TCCATTCACA CGAGGTCCTC CTCGCTCTGG CCGCTCTTGC TGCTCCTGTC	720
55	TGAAGAWATC AGACTGATTT CCTCTTAAGA CTCCTAGGGA TGTGGTGAAG AGCTGGGACT	780
	CAAGTGCAGT CCACGGTGTG AAACATGAGG GARGTGAGGT GTCCGTCCAC TTCCCCCATA	840

60

180

240

300

360

	CAGCCCCATC	TGGATGTGAG	GTGGGGTGGA	GACATCATGG	GGTGATTGCA	GAAAGGGGGA	960
•	GTGGCGGCCC	ACGCAGCTTC	TGCTGAGGAG	CTGACCGCTC	TGAGCTGTTC	TGTTTCGTAT	1020
5	TGCTGCTCTG	TGTCTGCATG	TATTGTGACC	GTGCGGCTCC	ACCTCTTCCA	GCTGCTGCTA	1080
	CAGCTGAGGC	CTGGATCCCG	GCCTTTCCCT	GTGACTTACG	TGTCTGTCAC	CGGCANGCAG	1140
10	CCCTACAAAT	CCTGGTGACC	TGCTCTCCCA	AGAACAGAGC	CTGTCCCCAG	ATGTCCCAGT	1200
	AGCGATGAGT	AACAGAGGTG	GCTGTGGACT	TCCTCTACTT	CTCCTTGCTG	GATCAGGGCC	1260
	TTCCTGCCTC	CCGCTGGGCA	GGTCTGGCCT	TGCTCTCTTG	GCAGGGCCCC	AGCCCCTCTG	1320
15	ACCACTCTGC	AGCTCACCAT	GCAGCTGATG	CCAAAGTTGT	GGTGTCCAGT	GTGCAGCAGC	1380
	CCTGGGAGCC	ACTGCCACCT	TCAGAGGGGT	TCCTTGCTGA	GACCCACATT	GCTTCACCTG	1440
20	GCCCCACCAT	GGCTGCTTGC	CTGGCCCAAC	CTAGCGTTCT	GTGCCATGCT	AGAGCTTGAG	1500
20	CTGTTGCTCT	TCTTCAGGGG	AGGAAATAGG	GTGGAGAGCG	GGAAGGGTCT	TGCTCCTAAG	1560
	TGTTGCTGCT	GTGGCTTTTT	TGCCTTCTCC	AAAGACGCAC	TGCCAGGTCC	CAAGCTTCAG	1620
25	ACTGCTGTGC	TTAGTAAGCA	AGTGAGAAGC	CTGGGGTTTG	GAGCCCACCT	ACTCTCTGGC	1680
	AGCATCAGCA	TCCTACTCCT	GGCAACATCA	GGCCAACGTC	CACCCCAGCC	TCACATTGCC	1740
30	AGATGTTGGC	AGAAGGGCTA	ATATTGACCG	TCTTGACTGG	CTGGAGCCTT	CAAAGCCACT	1800
	GGGATGTCCT	CCAGGCACCT	GGGTCCCATG	ACCAGCTCCC	CGTCTCCATA	GGGGTAGGCA	1860
	TTTCACTGGT	TTATGAAGCT	CGAGTTTCAT	TAAATATGTT	AAGAATCAAA	GCTGTCTTTG	1920
35	TTCAGGCTGC	TATAACAAAA	ATATAATAGC	CTGGGTGGCT	TAAAC		1965
•	<i>.</i>				•		
40	(2) INFORM	ATION FOR SE	EQ ID NO: 11	<b>L7</b> :			
	(i)	SEQUENCE CI	HARACTERIST:	ICS:			
			GTH: 503 ba E: nucleic a	-			
45		(C) STR	ANDEDNESS: OLOGY: line	double			
	(xi			: SEQ ID NO:	: 117:		
50		. ~		GCTGGAATTG		CCTCTGCACC	60
				CCACCATTCC			120
							120

TGCCTTTGGG AGAACCAGCC TCCTCCATGG AGGAAAGCTT GGGATCTGCC TTCCCACCTG

GGGAGGAGAG GGATCTGTGG AAAATCCTTC TGACGGACTT CCCCTCAGTG CCTGATCCAT

ACTCAATAGT AGAAAAAGTA AGAAATATAC AAAGATAGCA GATACACGGA GACAGTTCCC

CAAATAGCTG AGCGAWTAGC GCAGAAGCAA TATTGAAGAC CTAATAGCTG AGACATTTCC

	AGAACTGATA AAGTGCATCC AGCCACAGAT CAAGCAGCCC AGAAAATTCC AGGCAGCATC	420
5	AACAAATAAA TAGCCCCACA TGCACCCGTG AAAATGCAGA AGACCAAACA AAAAAGTCCG	480
	GTCAACAGCC AGAGTTAAAG AGG	503
10	(2) INFORMATION FOR SEQ ID NO: 118:	
	(i) SEQUENCE CHARACTERISTICS:	•
15	(A) LENGTH: 1133 base pairs (B) TYPE: nucleic acid	
10	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 118:	
	GGCACAGCTT GGAATGAACC CCTGTGGATA AGGGGGGACTA TTAGATAGAA TAAACATCAA	60
	TARATGCTTG ATGARTARAC GCTARTCCTA CCTTCCCAGC CTGACACCTC CCAGTGGACA	120
25	CCACACTTCA CTTGAAGCCT TAGAAACCTT TCCCACCCAT GCTTCCAGCC CTGGCTTCAT	180
	GTTGCCATTT CTCACCCCCA GAACAGGCCG CCCGCCTGAA GAAACTACAA GAGCAAGAGA	240
30	AACAACAGAA AGTGGAGTTT CGTAAAAGGA TGGAGAAGGA GGTGTCAGAT TTCATTCAAG	300
	ACAGTGGGCA GATCAAGAAA AAGTTTCAGC CAATGAACAA GATCGAGAGG AGCATACTAC	360
	ATGATGTGGT GGAAGTGGCT GGCCTGACAT CCTTCTCCTT TGGGGAAGAT GATGACTGTC	420
35	GCTATGTCAT GATCTTCAAA AAGGAGTTTG CACCCTCAGA TGAAGAGCTA GACTCTTACC	480
	GTCGTGGAGA GGAATGGGAC CCCCAGAAGG CTGAGGAGAA GCGGAACNIG AAGGAGCTGG	540
40	CCCAGAGGCA ANGAGGAGGA GGCAGCCCAG CAGGGGCCTG TGGTGGTGAG CCCTGCCAGC	600
	GACTACAAGG ACAAGTACAG CCACCTCATC GGCAAGGAG CAGCCAAAGA CGCAGCCCAC	660
	ATGCTACAGG CCAATAAGAC CTACGGCTGT KTGCCCGTGG CCAATAAGAG GGACACACGC	720
45	TCCATTGAAG AGGCTATGAA TGAGATCAGA GCCAAGAAGC GTCTGCGGCA GAGTGGGGAA	780
	GAGTTGCCGC CAACCTCCTA GGCGCCCCGC CCAGCTCCCT TTGACCCCTG GGGCAGGGCA	- 840
50	GGGGCAGGG AGAGACAAGG CTGCTGCTAT TAGAGCCCCAT CCTGGAGCCC CACCTCTGAA	900
	CCACCTCCTA CCAGCTGTCC CTCAGGCTGG GGGAAAACAG GTGTTTGATT TGTCACCGTT	960
	GGAGCTTGGA TATGTGCGTG GCATGTGTGT GTGTGTGTGA GAGTGTGAAT GCACAGGTGG	1020
55	GTATTTAATC TGTATTATTC CCCGTTCTTG GAATTTTCTT CCCATGGGGC TGGGGTACTT	1080
	TACATTCAAT AAATACTGTT TAACCCAAAA AAAAAAAAA AAAAGAAAGA AGN	1133

# (2) INFORMATION FOR SEQ ID NO: 119:

	(i) SEQUENCE CHARACTERISTICS:	
5	(A) LENGTH: 1101 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 119:	
	GGGCACAGCT GAAGCTGCAG ACCTCCCCAG GGGATGGCTC CTCTCCCCCA GGAGCCCCGA	60
15	GGCAGGGGAG GCAGAAAGCC TGGGCTCTGG GGGGTGGCCT GCGGACAGCT GTGCTGTGGG	120
	CCGGGGGCTG GGCCTGTCCC ACAGGGNCGT GGAGCTCGTG GTTCTGAGCA GCCAGCTGGG	180
	TGGTGTCTGG GGATAGCTGG GAGGCACAGC GGCTGCCATG TGGGACTGGG ACTGGAGTGC	240
20	TCCCTGGTCT TGGCCTCTGT GGCTCAGCCT TGCTCTGGTC TGCCTGAGTG CAGGGGCCAA	300
	GGGGCACAGG GCCAGTGAGG CCGGCCACGC TCGGGCCCTC ACCTGTGAGA TGGGGTCGGA	360
25	ATTTKACACA GCCTANGGCT TGGTTCTTGG TKGTNGAMCG TGGACTYCTK AGAACGGGAG	420
	TGCTGGTCCT GAAAGGCGTG GTTGGAGACC AGCTGCTTTT CTCGCTGTTT TTCTCTTAGG	480
	AGATTAAACA AAAACAGAAA GCACAAGACG AACTCAGTAG CAGACCCCAG ACTCTCCCCT	540
30	TGCCAGACGT GGTTCCAGAC GGGGAGACGC ACCTCGTCCA GAACGGGATT CAGCTGCTCA	600
	ACGGGCATGC GCCGGGGGCC GTCCCAAACC TCGCAGGGCT CCAGCAGGCC AACCGGCACC	660
35	ACGGACTCCT GGGTGGCGCC CTGGCGAACT TGTTTGTGAT AGTTGGGTTT GCAGCCTTTG	720
	CTTACACGGT CAAGTACGTG CTGAGGAGCA TCGCGCAGGA GTGAGGCCCCA GGCGCCGAGA	780
10	CCCAAGGCGC CACTGAGGGC ACCGCGCACC AGAGCGTGAC CTCGGCAGGC TGGACACACT	840
10	GCCCAGCACA GGCAGACCCA CCAGGCTCCT AGGTTTAGCT TTTAAAAACC TGAAAGGGGA	900
	AGCAAAAACC AAAATGTGTG ACTGGGCTTT GGAGGAGACT GGAGCCTCAG CCCTGTCCTG	960
15	GCCACGGGCC GCTGGGGCTG GTGTGGGTGG GCCTTGTGTG CTGGATTTGT AGCTTATCTT	1020
	CCGTGTTGTC TTTGGACCTG TTTTAGTAAA CCCGTTTTTC ATTTTAAAAA AAAAAAAAAA	1080

50

# (2) INFORMATION FOR SEQ ID NO: 120:

55 (i) SEQUENCE CHARACTERISTICS:

AAACTTTGGG GGGGGGCCCC N

(A) LENGTH: 282 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 120:	
	AGCTTCTCTG TCCAGTCTTG AACTCTGGGS TCTCTTGGAA CTTTCCTCAC CCCTCTCAGC	6
5	CTGAATATTC CTTCCATGGA TTCCACTCAA CCAGACTTTG GATCTGTGCC TACTTAATCA	120
	ACCITATETT TECAATATET TEGGGCCCAC CITCCACTCC TIGGITCITG TICCTCCTIG	180
10	GCCTAACTTG TCCCTTCTCC ACTTCACATC CCCGGTGGGA CAGCATTCCT CCTTCCTCCC	. 240
10	AACCTCCCTC CGTCTCARAA AAAAAAAAAA AAAAAAAAAA TT	282
15	(2) INFORMATION FOR SEQ ID NO: 121:	
20	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2635 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 121:	
20	TAAGGGGGTG TGTGCTCACC TCCTCCTGAC CCTTAACACT CCTGTCCTGC CCAGACCAAC	60
	AGAGAGAGCT GTCCCTGAGA CCCCGGAGAG AAGCAGCTGC CGAAAGCTGC AGCCTTTCCG	120
30	CACTCTGAGA CCATGATCTT CCTCCTGCCA GGGGAGAGCC ACCCACAGGC CATGTCCAGC	180
	CCCACTTCCC TCAGCCCCCA GGGYTTCCTT CTGGCCCCTC TGAGGATTCC CTAGGGCTGC	240
35	CCCGCAGAGG GGYTTCCCCA AGCTCTGTTT TGAAGCCTGC AATGTGGAAA AGTGAGAAGT	300
	CAGAGGGAAC AGGACAGGTG CAGCCGGGCT CTGAGGCCAC ACCTCACACC TCGCTGTTCC	360
	CCAACATCCC CTGAGCAGTG TGAGCTCATC TCACCAGATG AGAAGAGGCC CTGTGCATTT	420
40	YTTTTGTTG TTTGTTGCTG TTTTCCCCCA CCCATCCAGT TCTCCTCAGC AAAGCAAATT	480
	CCTTAACACC TTTGGTGGAG AATTTCTTAC CCAGACTTGG GGCTGTGATG CCCTTCAGTG	540
45	CGTGGTGAGT GCAGCGTGTG TGCGTGTGCC TGTGTGTGAA CCTGGGGGCC ATCCTGGTGG	. 600
	CCTGGGAGCG TGAGGAGAGG CCCCCTGTGT GCTGGGTGAG TGGTGGGTGT GGGGTCAATG	660
	CAGTGAGGCT CTCTGGGTGA GGCTCCCAAC CTGGCAGTCC CCAGCCTCCC AGCATCTGTG	720
50	AGCGTCTGTT GGACTTTACA GAAGAGCCTC ATCCYGTCTG CCCCTCACTC TGCCCTGGAA	780
	TCAACATCTT CCGAGTCCTT CTTGGGGGAA ATAGCAGAGC CCCACTTAAC TCCATAAACT	840
55	GCTTCCCATT CCGCAGCCCA GTTCTGATTG TTGAGGTGTC GCGTCGTTCC AGGTCCCCCA	900
	GTCCCCTCTT TCTCCTGTCC TCTCTCTGTC CTTCACCTCC CCACTCCAGC CCCGGCTCAG	960
	TTCAGGGAAA TGCTGTTCCA YATCAGCCCT CTGCTCTCTG AGGCAGCCGC GCCTCTGACT	1020
60	CGGAGCTACT TGAAACTTCT GCTCTTGCTA GGATTGGAGT CTACCTATCT CTTCCATTTG	1080

	TCCCAGCTGG	AGTTCTGGAA	CTTTCCTCCT	CGGGGTGGGG	GIGGGGGTT	TTAAGGATGC	114
5	TGGGGGGCCT	GGGGAAGGAA	. GGAGTTCAGA	GGAAGGGTGT	, cccararca	CTTSATGTCA	120
	CCCTCCGCTC	CTGGGACACG	TGCTCTCTCT	GTCTCTGGGT	CITCIGGGTG	TGCACGTTTG	126
	TGTGTCCTTG	TAAATATGTT	TTAGGAAGAA	AGCAAAAGGG	ACTGAACTAC	COTOTGGTAG	1320
10	GATTGCAGGG	GTCCAGCCTT	GCCTGTTTCC	GAAGCCCCCA	. CACTGCCTT	CGCCCCACTG	1386
	AGACTGGTCC	CCTCAAAAGG	TAGACAAAAC	AGCAGCTCCC	TGTGGAGCTG	AAGGCGGCC	1440
15	TCAAAGTGGC	TTTTTGTTAG	ACAAGGTTAA	GGTTTCCTCA	TGAGCAAGGI	TGCAGATCGG	1500
	TCCTTCCTCA	GCTCCTTGAT	TTGTGACCTT	. GYCCYYGGGG	CCTGCCAECC	AGCCCCTCCA	1560
	GTGCCCTCTC	CTCGATGCCT	CGCTCCTTCC	TGCCCCCACT	CCCCTGGETT	AGGEAGGTAG	1620
20	GGGAATTAGG	GCCATGCTGG	AAGAAGCTTA	ACCATGTGTT	CAAAGAACGG	TTTCTTGCTT	1680
	GCTTGGTCCT	GGAACTCCCC	TTGGCTGCCC	CAGGCCTCCT	TGGCCCATGG	GIGTIGGGG	1740
25	AGGTGGATGT	CAGATCTGGT	AGGTTGCAGC	AGAGAAAATA	AATGTGCTTT	GAGAGACCAC	1800
	TCAGAGAGGG	TCCAAGGGTG	ATGGAGAAGG	AAGCATGGCC	TGGGAGCTTG	GAAGGGARGG	1860
	GTGGTGGGTG	GCGGCATCTT	GACTGCCCCC	TGTTGTCCCA	CACGTGGGGG	GTGGTCACCC	1920
30	CYCTTCACTC	CAGCCCGCCT	GCCTTCAGCC	TTCCATGAGC	TTCACCTGCT	TCCAACTTCA	1980
	CTTTGGAGGG	GGTGGGGTCC	GTTGGCATCA	ACACGGGGAC	CCTCTGCTTC	ACCAAAGCCC	2040
35	GAGCCCTCAG	CCCCTGGGGA	GAACAAATGG	CTGAGCTTTG	ATACCTGGGG	TCGTCGAGAG	2100
	GCTGCGGGCT	GGCGGCAGTC	CCAGGGGAGA	GACACCACAG	AAGGAGACCC	AGACATOCCG	2160
	AGGAAGTTCC	CAGCAGAGCA	AACTGCTTTC	CAGCCTGAAG	CCTGCTTAAA	CTGTGTGATG	2220
40	TGCAATAACT	GAGCTTAGAG	TTAGGAATTG	TGTTCAAGTG	CTTTGGATTTC	CGTCTGTAGA	2280
	TTTAACTGCT	GAAATTGTAT	CTCTCAGTAA	TTTTAGATGT	CTTTTAAAAA	ATTGAAAAAC	2340
<b>1</b> 5	AAAGTGTTAG	ACTGTGTGCG	TGTGCGTTGA	TGGGCACTCA	AGAGTCCCFT	GASTCATCCA	2400
	GCCCTGCCTT	TCCCCTGCGC	CCCCATCCTC	TCACGTCCCG	ccasecatcc	ACTTGGGGAC	2460
	CCTGCCTCGT	GTCGTCTTTA	TCTGCCTATT	ACTCAGCCTA	AGGAAACLAG	TACACTCCAC	2520
50	ACATGCATAA	AGGAAATCAA	ATGTTATTTT	TAAGAAAATG	GAAAATAAA	ACTITATAAA	2580
	CACCAAAAAA	АААААААА	ACCONGGGGG	GGGCCGGTA	ACCCATTTCG	CCTAA	2635

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 994 base pairs

<sup>(2)</sup> INFORMATION FOR SEQ ID NO: 122:

(B) TYPE: nucleic acid

	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 122:	
	GAATTCGGCA GAGGTTCGGC GAAGATAGGG AATAAGGAAG CACAGGAGTA GGGGAGAAGG	60
10	AAGCACAGGA GTAGGGGAGA TATACAGCGG TCAGGATAAG GGGGAAAGGG CGGTGGTTGC	120
10	SCAAGAGGTG AAACAAGATG TGAGAGACAA GGGGTAGGGA AGAAATGGGG CAGCGGTTAG	180
	GTTCAGAAGC GCATAGACCG TGGCGGACGG GCAATGCGAG GGGCACAGAA AGGAACTGAG	240
.15	GGGTGGGCTA TTTTAARGGA GATGGTCCTT CAGCCCTCTT YTTTTCTGCG TAGTTCTCCT	300
	CCTCCAGGCC GCGCGCGGAT ATGTCGTCCG GAAACCAGCC CAGTCTAGGC TGGATGATGA	360
20	CCCACCTCCT TCTACGCTGC TCAAAGACTA CCAGAATGTC CCTGGAATTG AGAAGGTTGA	420
20	TGATGTCGTG AAAAGACTCT TGTCTTTGGA AATGGCCAAC AAGAAGGAGA TGCTAAAAAT	480
	CAAGCAAGAA CAGTTTATGA AGAAGATTGT TGCAAACCCA GAGGACACCA GATCCCTGGA	540
25	GGCTCGAATT ATTGCCTTGT CTGTCAAGAT CCGCAGTTAT GAAGAACACT TGGAGAAACA	600
	TCGAAAGGAC AAAGCCCACA AACGCTATCT GCTAATGAGC ATTGACCAGA GGAAAAAGAT	660
30	GCTCAAAAAC CTCCGTAACA CCAACTATGA TGTCTTTGAG AAGATATGCT GGGGGCTGGG	720
50	AATTGAGTAC ACCTTCCCCC CTCTGTATTA CCGAAGAGCC CACCGCCGAT TCGTGACCAA	780
	GAAGGCTCTG TGCATTCGGG TTTTCCAGGA GACTCAAAAG CTGAAGAAGC GAAGAAGACC	840
35	CTTAAAGGCT GCAGCAGCAG CCCAAAAACA AGCAAAGCGG AGGAACCCAG ACAGCCCTGC	900
	CAAAGCCATA CCAAAGACAC TCAAAGACAG CCAATAAATT CTGTTCAATC ATTTAAAAAA	960
40	AAAAAAAAA AAAAAAAAA AAAAAGGGGA GGGG	994
	(2) INFORMATION FOR SEQ ID NO: 123:	
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1542 base pairs  (B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 123:	
55	GGCASAGCCA CCTCGGCCCC GGGCTCCGAA GCGGCTCGGG GGCGCCCTTT CGGTCAACAT	60
	CGTAGTCCAC CCCCTCCCCA TCCCCAGCCC CCGGGGATTC AGGCTCGCCA GCGCCCAGCC	120
60	AGGGAGCCGG CCGGGAAGCG CGATGGGGGC CCCAGCCGCC TCGCTCCTGC TCCTGCTCCT	180
	GCTGTTCGCC TGCTGCTGGG CGCCCGGCGG GGCCAACCTC TCCCAGGACG ACAGCCAGCC	240

GCTGTTCGCC TGCTGCTGGG CGCCCGGCGG GGCCAACCTC TCCCAGGACG ACAGCCAGCC

	CTGGACATCT GATGAAACAG TGGTGGCTGG TGGCACCGTG GTGCTCAAGT GCCAAGTGAA	300
5	AGATCACGAG GACTCATCCC TGCAATGGTC TTAACCCTGC TCAGCAGACT CTCTACTTTG	360
J	GGGAGAAGAG AGCCCTTCGA GATAATCGAA TTCAGCTGGT TAMCTCTACG CCCCACGAGC	420
	TCAGCATCAG CATCAGCAAT GTGGCCCTGG CAGACGAGGG CGAGTACACC TGCTCAATCT	480
10	TCACTATGCC TGTGCGAACT GCCAAGTCCC TCGTCACTGT GCTAGGAATT CCACAGAAGC	540
	CCATCATCAC TGGTTATAAA TCTTCATTAC GGGAAAAAGA CACAGCCACC CTAAACTGTC	600
15	AGICTICIGG GAGCAAGCCT GCAGCCCGGC TCACCTGGAG AAAGGGTGAC CAAGAACTCC	.660
	ACGGAGAACC AACCCGCATA CAGGAAGATC CCAATGGTAA AACCTTCACT GTCAGCAGCT	720
	CGGTGACATT CCAGGTTACC CGGGAGGATG ATGGGGGCGAG CATCGTGTGC TCTGTGAACC	780
20	ATGAATCTCT AAAGGGAGCT GACAGATCCA CCTCTCAACG CATTGAAGTT TTATACACAC	840
	CAACTGCGAT GATTAGGCCA GACCCTCCCC ATCCTCGTGA GGGCCAGAAG CTGTTGCTAC	900
25	ACTGTGAGGG TCGCGGCAAT CCAGTCCCCC AGCAGTACCT ATGGGAGAAG GAGGGCAGTG	960
<del>-</del>	TGCCACCCCT GAAGATGACC CAGGAGAGTG CCCTGATCTT CCCTTTCCTC AACAAGAGTG	1020
	ACAGTGGCAC CTACGGCTGC ACAGCCACCA GCAACATGGG CAGCTACAAG GCCTACTACA	1080
30	CCCTCAATGT TAATGACCCC AGTCCGGTGC CCTCCTCCTC CAGCACCTAC CACGCCATCA	1140
	TCGGTGGGAT CGTGGCTTTC ATTGTCTTCC TGCTGCTCAT CATGCTCATC TTCCTTGGCC	1200
35	ACTACTTGAT CCGGCACAAA GGAACCTACC TGACACATGA GGCAAAAGGC TCCGACGATG	1260
	CTCCAGACGC GGACACGGCC ATCATCAATG CAGAAGGCGG GCAGTCAGGA GGGGACGACA	1320
	AGAAGGAATA TTTCATCTAG AGGCGCCTGC CCACTTCCTG CGCCCCCCAG GGCCCTGTGG	1380
40	GGACTTGCTG GGGCCGTCAC CAACCCGGAC TTGTACAGAG CAACCGCAGG GGCCGSCCCT	1440
	CCCGNTTGTT CCCCAGCCCA CCCACCCCCT TGTTACAGAA TGTYTKGTTT GGGGTGCGGT	1500
45	TITGIWATTG GITTNGGATN GGGGAAGGGA GGGANGGCGG GG	1542
	(2) INFORMATION FOR SEQ ID NO: 124:	
50	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1390 base pairs (B) TYPE: nucleic acid	
55	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	•
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 124:	

CAAGCTCTAA TACGACTCAC TATAGGGAAA GCTGGTACGC CTGCAGGTAC CGGTCCGGAA

	TTCCCGGGTC	GACCCACGCG	TCCGGGCCTC	AGGGTGGACG	CATGGTTCTG	CACTGAGGCC	12
	CTCGTCATGG	TGGCGCCTGT	GTGGTACTTG	GTAGCGGCGG	CTCTGCTAGT	CGGCTTTATC	180
5	CTCTTCCTGA	CTCGCAGCCG	GGGCCGGGCG	GCATCAGCCG	GCCAAGAGCC	ACTGCACAAT	240
	GAGGAGCTGG	CAGGAGCAGG	CCGGGTGGCC	CAGCCTGGGC	CCCTGGAGCC	TGAGGAGCCG	300
10	AGAGCTGGAG	GCAGGCCTCG	GCGCCGGAGG	GACCTGGGCA	GCCGCCTACA	GGCCCAGCGT	360
10	CGAGCCCAGC	GGGTGGCCTG	GGCAGAAGCA	GATGAGAACG	AGGAGGAAGC	TGTCATCCTA	420
•	GCCCAGGAGG	AGGAAGGTGT	CGAGAAGCCA	GCGGAAAYTC	ACCTGTCGGG	GAAAATTGGA	480
15	GCTAAGAAAC	TGCGGAANNT	GGAGGAGAAA	CAAGCGCGAA	AGGCCCAGCK	TGAGGCAGAG	540
	GAGGCTGAAC	GTGARGWGCG	GAAACGACTC	GAGTCCCAGC	GCGAATGAGT	GGAAGAAGGA	600
20	GGAGGAGCGG	CTTCGCCTGG	AGGAGGAGCA	GAAGGAGGAG	GAGGAGAGGA	AGGCCCGCGA	660
20	GGAGCAGGCC	CAGCGGGAGC	ATGAGGAGTA	CCTGAAACTG	AAGGAGGCCT	TTGTGGTGGA	720
	GGAGGAAGGC	GTAGGAGAGA	CCATGACTGA	GGAACAGTCC	CAGAGCTTCC	TGACAGAGTT	780
25	CATCAACTAC	ATCAAGCAGT	CCAAGGTTGT	GCTCTTGGAA	GACCTGGCTT	CCCAGGTGGG	840
	CCTACGCACT	CAGGACACCA	TAAATCGCAT	CCAGGACCTG	CTGGCTGAGG	GGACTATAAC	900
30	AGGTGTGATT	GACGACCGGG	GCAAGTTCAT	CTACATAACC	CCAGAGGAAC	TGGCCGCCGT	960
50	GGCCAACTTC	ATCCGACAGC	GGGCCGGGT	GTCCATCGCC	GAGCTTGCCC	AAGCCAGCAA	1020
	CTCCCTCATC	GCCTGGGGCC	GGGAGTCCCC	TGCCCAAGCC	CCAGCCTGAC	CCCAGTCCTT	1080
35	CCCTCTTGGA	CTCAGAGTTG	GTGTGGCCTA	CCTGGCTATA	CATCTTCATC	CCTCCCCACC	1140
	ATCCTGGGGA	AGTGATGGTG	TGGÇCAGGCA	GTTATAGATT	AAAGGCCTGT	GAGTACTGCT	1200
40	GAGCTTGGTG	TGGCTTGGTG	TGGCAGAAGG	CCTGGCCTAG	GATCCTAGAT	AAGCAGGTGA	1260
	AATTTAGGCT	TCAGAATATA	TCCGAGAGGT	GGGGAGGGTC	CCTTGGAAGC	TGGTGAAGTC	1320
	CTGTTCTTAT	TATGAATCCA	TTCATTCAAG	AAAATAGCCT	GTTGCAAAAA	AAAAAAAA	1380
45	AAAAACTCGA						1390

# 50 (2) INFORMATION FOR SEQ ID NO: 125:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1288 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 125:

60 GGCGCGGGG TGAAAGGCGC ATTGATGCAG CCTGCGGCGG CCTCGGAGCG CGGCGGASCA

WO 98/54963 PCT/US98/11422

378

	GACGCTGACC ACGTTCCTCT CCTCGGTCTC CTCCGCCTCC AGCTCCGCGC TGCCCGGCAG	120
_	CCGGGAGCCA TGCGACCCCA GGGCCCCGCC GCCTCCCGC AGCGGCTCCG CGGCCTCCTG	180
5	CTGCTCCTGC TGCTGCAGCT GCCCGCGCCG TCGAGCGCCT CTGAGATCCC CAAGGGGAAG	240
	CAAAAGGCGC ATCCGGCAGA GGGAGGTGGT GGACCTGTAT AATGGAATGT GCTTACAAGG	. 300
10	GCCAGCAGGA GTGCCTGGTC GAGACGGGAG CCCTGGGGCC AATGGCATTC CGGGTACACC	360
•	TGGGATCCCA GGTCGGGATG GATTCAAAGG AGAAAAGGGG GAATGTCTGA GGGAAAGCTT	420
15	TGAGGAGTCC TGGACACCCA ACTACAAGCA GTGTTCATGG AGTTCATTGA ATTATGGCAT	480
13	AGATCTTGGG AAAATTGCGG AGTGTACATT TACAAAGATG CGTTCAAATA GTGCTCTAAG	540
	AGTTTTGTTC AGTGGCTCAC TTCGGCTAAA ATGCAGAAAT GCATGCTGTC AGCGTTGGTA	600
20	TTTCACATTC AATGGAGCTG AATGTTCAGG ACCTCTTCCC ATTGAAGCTA TAATTTATTT	660
	GGACCAAGGA AGCCCTGAAA TGAATTCAAC AATTAATATT CATCGCACTT CTTCTGTGGA	720
25	AGGACTTTGT GAAGGAATTG GTGCTGGATT AGTGGATGTT GCTATCTGGG TTGGCACTTG	780
23	TTCAGATTAC CCAAAAGGAG ATGCTTCTAC TGGATGGAAT TCAGTTTCTC GCATCATTAT	840
	TGAAGAACTA CCAAAATAAA TGCTTTAATT TTCATTTGCT ACCTCTTTTT TTATTATGCC	900
30	TTGGAATGGT TCACTTAAAT GACATTTTAA ATAAGTTTAT GTATACATCT GAATGAAAAG	960
	CAAAGCTAAA TATGTTTACA GACCAAAGTG TGATTTCACA TGTTTTTAAA TCTAGCATTA	1020
35	TTCATTTTGC TTCAATCAAA AGTGGTTTCA ATATTTTTTT TAGTTGGTTA GAATACTTTC	1080
33	TTCATAGTCA CATTCTCTCA ACCTATAATT TGGGAATATT GTTGTGGTCT TTTGTTTTTT	1140
	CTCTTAGTAT AGCATTTTTA AAAAAATATA AAAGCTACCA ATCTTTGTAC AATTTGTAAA	1200
40	TGTTAAGAAT TTTTTTTATA TCTGTTAAAT AAAAATTATT TCCMACAACC TTAAAAAAAA	1260
	AANAAAAA AAAAAAAAA	1288
45		
	(2) INFORMATION FOR SEQ ID NO: 126:	
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1517 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
55	(WILL CHARMON DESCRIPTION) SEC ID NO. 126:	

AGTGGCTTAA AGGCATCGTT TTAGGGATTA CTGGGAAGTA TCTTCAAAGT AATACATGAG

AAACATTCCT TCCTAAATCC TTTATTATAT TGAATATCGT ATTAATTGGT TTTCAGAGGT

60

60

	TAAATTAACC	ATGTATTCCT	GCAATAAATG	TCACTTGTNT	CTTGTATATA	ATCTTTTTTA	180
	TATATTACCG	GATTGATTCA	TTAGTATTTT	GTTGAGGATT	TTTGTGTCTA	TATTCATAAG	240
5	AGATGCTGGT	CTGCAGTTTT	CTTTTTTTGT	GATAATCTGG	TTTTTGTATC	AGTAATACAG	300
	GCCCCATGAA	ACGAGTTGGG	AAGTGTTCAC	CTCTCTTGTA	TTTTTTCAAG	AGTTTGTGAA	360
10	GAATTGCTAT	TAATTCTTTA	AATGTTTGGT	AGAATCTACC	ATTGAAATCA	TCTCTCCTGG	420
10	GCTTTTTTT	GAGGGAAGTG	TTCTGATAAC	TAATTCAGTA	TCTACTTTTT	ATAGCTCTGT	480
	TCAGATTTTG	CTTCTTCCTG	AGTTAGTTTT	GGTAATTTGT	GTATCTCTAG	GARTTTGTCC	540
15	ATTTCATTTA	TCTCATTTGT	TGGCATAAAT	TAAACTAAAT	TTGGCCTGAG	CCTACCTGTA	600
	TATCTTGAGT	CCCTCTGTAA	GGAACTGTAG	CCTAACTTGT	ACATAAACAA	ACTGAAATCC	660
20	TAAATTAGGA	ATGTAGTTTT	TGTAACAGCT	CCTGAGTCTC	AGGCAGTCAC	AGCAGYCAAG	720
20	TCTGTCAATT	GCAGGCTGCT	AACTAAGCAG	CCCATGSTCA	AATGAGGCAA	AAACCTTTGC	780
	TTTTAACACA	TAGTATAGCT	TTGTAATCCT	TTTCTTGCAC	ACTCGGGTAA	TTTCTTCCTT	840
25	TTTCATTCCC	KGWATTTTCC	AKGAATATGA	RTCTYCCTTT	TTTCCCCTCC	TGTCAGTCTA	900
	GCTAATGGTT	TGTCAATTTT	GTTGATCTTT	TGAARAACAA	ACCTTTGGTT	CCACTTTCTT	960
30	GTTGCATATG	CTGARTATTC	TCATAATTGG	AGTGGAAAGC	TGATCTTTGA	TTACTTATTT	1020
50	TACTTAGGGC	TGAGGAGTTC	ATGGACTTCG	CAAAACCTCC	TTGAATCTAA	ATTGCATCTT	1080
	CTTTCCTGGT	TTCTGGGCTG	AAACATGTTT	TTTCCCATCT	WANAWACCCI	TGGTCTTTTC	1140
35	ATKGGCGATT	AAGACTAGAG	AAAGTTCTAG	ATMCCTTGTC	CTTTTATGCT	GTCATTTTGT	1200
	TTAAAGGCTT	TCTATGTAGI	AAAACTATCI	ATATAGACAA	AATAGAGCCT	TGAGTTGTGG	1260
40	TCTTGAATTT	GATCAACATG	ATTTACCACA	TTCTGTACTC	GATATTTCTT	CACCTGCTGC	1320
40	TACTGTAAAC	CATTTTATTO	TTGGATCTTC	TGTAGAGTAT	ATTATCACAC	GTACTTTTTA	1380
	CAGGGGTGTC	TAATCTTTTC	GCTTCCCTGC	GCACATTGA	AGAAGAAGA	A TIGTCTIGGG	1440
45	CCACACATCA	A AATACGCTA	CACTAATAA	r agttgatgac	CTAAAAAAA	A AAAAAAAAAG	1500
	GCAAAAAAGI	N CCCAAAA					1517

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# (2) INFORMATION FOR SEQ ID NO: 127:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1073 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 127:

	TGAATCTATT CTTTGAACAT TCTACAACAA GAATTACATT ATACTGTTAT ACCAGAGTAC	60
5	TTCTGCAGTG TGAAATAGAT TGGTTTGGAA AATGAACCTG GCTTTGCTAT AAATTACATT	120
J	CACAGGCCTT TTTGCAAATG TGTAACTTGC CTATCAAAGT AGTTTGTAGG GCAAATGCAG	180
	AATATATGTC TCCATCTGGT AAAGTACCTT WTAYTCATGT GGGAAATCAA GTAGTATCAG	240
10	AACTTGGTCC AATAGTCCAA TTTGTTAAAG CCAAGGGCCA TTCTCTTAGT GATGGGCTGG	300
	AGGAAGTCCA AAAAGCAGAA ATGAAAGCTT ACATGGAATT AGTCAACAAT ATGCTGTTGA	360
1 5	CTGCAGAGCT GTATCTTCAG TGGTGTGATG AAGCTACAGT AGGGRMGATC ACTCATGMTA	420
15	GGTATGGWTC TCCTTACCCT TGGCCTCTGW WTCATATTTT GGCCTATCAA AAACAGTGGG	480
	AAGTCAAACG TAAGNTGAAA GCTATTGGAT GGGGAAAGAA GACTCTGGAC CAGGTCTTAG	540
20	AGGATGTAGA CCAGTGCTGT CAAGCTCTCT CTCAAAGACT GGGAACACAA CCGTATTTCT	600
	TCAATAAGCA GCCTACTGAA CTTGACGCAC TGGTATTTGG CCATCTATAC ACCATTCTTA	660
25	CCACACAATT GACAAATGAT GAACTTTCTG AGAAGGTGAA AAACTATAGC AACCTCCTTG	720
25	CTTTCTGTAG GAGAATTGAA CAGCACTATT TTGAAGATCG TGGTAAAGGC AGGCTGTCAT	780
	AGAGTTATGT GTTAGTCTCA GGAGTCTTAA CTTTTGAAAT ATGTTTACT TGAATGTTAC	. 840
30	ATTAGATATT GGTGTCAGAA TTTTAAAACC AAATTACTGC TTTTTGAAAC CTCAAATTAT	900
	ATAATGTATC TTATGTATGT GCTTTATATT GTTATTTGTG TATACATTAA AATAATTCTG	960
25	AATTATTTAA TCTGATATGT TGTATTCTGT ATCTTGAAAT TTTTGTTTCC TTGAAACATG	1020
35	CATGCATTTA AAAATAAAGC TTAAACAACT GTAAAAAAAAA AAAAAAAAAA	1073
		-
40		
	(2) INFORMATION FOR SEQ ID NO: 128:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 300 base pairs	
45	<ul><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li></ul>	
	(D) TOPOLOGY: linear	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 128:	
	CAACCCCTGC CTTTTTTTTG TTTTCCATTT GCTTGGTAGA TCTTCCTCCA TCCCTTTATT	60
	TTGAGCCTAT GTGTGTCTCT GCCCGTGAGA TGAGTCTCCT GAATACAGCA CACTTACTGG	120
55	TCTTGACTCT GTATCCAATT TGCCAGTCTG TGTCTTTCAT TTGGAGCATT TAGCCCATTT	180
	ACATTTAAGG TKAATATTGT TATGTGTGAA TTTRATCYTR TCATTATGWT GTTAGCTGGT	240
60	TATTTTGCTT GTTAGTTGAT GCAGTTTCTT CCNGGCATCA ATGGTCTTTA CAANTTGGCA	300
50		

(2) INFORMATION FOR SEQ ID NO: 129:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1275 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

10 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 129:

	(111)	
15	GGCAGAGCCT GTCCCTGCTG CCCCTGCAAA AAAAACCCCC TCTGGTGTGA GCAGGATGGT	60
13	TEGAGGITAT GIGAGCICCI TCTCCTITCC TCCAGITTCC TCTTCCCTTC TCCTCCCTGC	<sub>-</sub> 120
	CTCTTTTGCT TTTCCCTTTC TTCCTGGTAC CCCCTGCCCA TTCCTGTATT TTCTCCCATC	180
20	GCCATTCTCC CCTCTCCCAC TGTCCCTAAC CCGTTCAAAC TCTTTCCTCT TAAATGGTTG	240
	AGATTTTCTC TCACCAAGCA CACCCCAGTA TTAATTAAAC TAGCTGCAAA CAGGCAGCAA	300
25	GTGGTCTACC ATGACAGATG GGTTTTGTGT GTGTGTGTGT GTGTGTAATT GTAATAAAAC	360
20	ATATTGARTC ACTCAATAAA CACAGAGTGT CTACTACATG TATCARGCAC TATCATAGAT	420
	GCTAATTAAC GAAACTGAAA TGGCCAGGCC CTCACAGTGG CTCATGCCTA TAATCCCAGC	480
30	ACTITIGGAG GATGAGGCAG GAGGATCACT TGAGGCCGGG AGTTCAAGAC CAGCCTGGGC	540
,	AACATAGTAA GACTCCATCT CTACAAAAAA AAAATTTTTT TTATTATACT TTAAGTTTTG	600
35	GGTTACATGT GCAGAACGTG TAGTTTTGTT ACATAGGTAT ATACGTCCCC TGGTAGTTTG	660
	CTGCACCCAT CAACCCATCA CCTACATTAG GTATTTCTCC TAATGTTACC CCTCTCCTAG	720
	CCCCCCACCC CGTGACAGGC CCTGGTGTGT GATGTTCCCC TCCCTGTGTC CATGTGTTCT	780
40	CATTGGTCAA CTCTCACCTA TGGAGTGAGA ACATGTGGTA TTTGGTTTTC TGATCTTGTG	840
	ATAGCTTGCT GAGAATGTKG GTTTCCAGCT TTATCCACGT CCCTGCAAAG GGCATAAACT	900
45	CATCCCTTTT TATGGCTGCA TAGTGTTCCA TGGTGTATAC GTGCCACATT TTCTTAATCT	960
	ATCATTGATG GACAAGTTTT GCTATTGTGA ATAGTGCCAC AATAAACATA CGTGTGCGTG	1020
	TGTCTTTATA GCAGCATGAT TTATAATCCT TTGGGTATAT ACCCAGTAAT GGGATCACTG	1080
50	AGTCAAATGG TATTTCTCGT TCTAGATCCG TAAGGAATTG CCACACTGTC TTCCACAATG	1140
	TTTGAACTAA TNTACACTCC CACCAACAGT GTAAAAGTGT TTCTATTTTT CCACAACCTC	1200
55	TCCAACATCT GTTATTTCCT GACTTTTTAA TGAACGTCAT TCTAACTGGC GTGAGATGGT	1260
-	ATCTCATTGT GGTTT	1275

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(2)	INFORMATION	FOR	SEQ	ID	NO:	130:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 472 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 130:

10 CNGAAACCCC GTGAACCCTC CCCGGGTTAA AAAGCCCCCC CTAAATGGGG GGAACGCYTC 60 ACACGTTATA AAAAAGCACT AGAATGTTTT GAAAGCGAGA AACAACAGCT GTGTAGGGTA 120 GCTAGCAGTT AGTGTTGTAC AGAAGACAGA TATTTGTGCA TTTYTGCATT TTCTAAGTTT 180 15 GCTGCAATGA GCATGTATTA CTTTCATAGT TATAAAACAC ATGCAAAATG CCCTTTTAAA 240 ATGAAAAAA ATCCATGAGT GTAAGTGATA TATATGCTTT GGAAAGCCTG GGACGGTCAT 300 20 TGTTTACTCT CAATAGTATG TGTTTGCCTT TGTCTTTTTG AGACATTTTG TTTTAATCTG 360 TTGATGACAA TAACCTGTTG ATAATATAAC TTGATAACAA ATAAAATGAC TTATGATTGA 420 472 25 

#### (2) INFORMATION FOR SEQ ID NO: 131:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1950 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

#### (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 131:

ACCTCTCAGA ATCTTCTCTC AGCAACCTGA GTCTTCGCCG TTCCTCAGAG CGCCTCAGTG 60 . 40 120 ACACCCCTGG ATCCTTCCAG TCACCTTCCC TGGAAATTCT GCTGTCCAGC TGCTCCCTGT 180 GCCGTGCCTG TNATTCGCTG GTGTATGATG AGGAAATCAT GGCTGGCTGG GCACCTGATG 45 ACTCTAACCT CAACACAACC TGCCCCTTCT GCGCCTGCCC CTTTNTGCCC CTGCTCAGTG 240 TCCAGACCNT TGATTCCCGG CCCAGTGTCC CCAGCCCCAA ATCTGCTGGT GCCAGTGGCA 300 [ GCAAAGATGC TCCTGTCCCT GGTGGTCCTG GCCCTGTGCT CAGTGACCGA AGCTCTGCCT 50 TGCTCTGGAT GAGCCCCAGC TCTGCAACGG GCACATGGGG GGAGCCTCCC GGCGGGTTGA 420 GAGTGGGGCA TGGGCATACC TGAGCCCCCT GGTGCTGCGT AAGGAGCTGG AGTCGCTGGT 480 55 AGAGAACGAG GGCAGTGAGG TGCTGGCGTT GCCTGAACTG CCCTCTGCCC ACCCCATCAT 540 CTTCTGGAAC CTTTTGTGGT ATTTCCAACG GCTACGNCTG CCCAGTATTC TACCAGGCCT 600 GGTGCTGGCC TCCTGTGATG GGCCTTCGMA CTCCCAGGCC CCATCTCCTT GGCTAACCCC 660 60

	TGATCCAGCC	TCTGTTCAGG	TACGGCTGCT	GTGGGATGTA	CTGACCCCTG	ACCCCAATAG	720
5	CTGCCCACCT	CTCTATGTGC	TCTGGAGGGT	CCACAGCCAG	ATCCCCCAGC	GGGTGGTATG	780
J	GCCAGGCCCT	GTACCTGCAT	CCCTTAGTTT	GGCACTGTTG	GAGTCAGTGC	TGCGCCATGT	840
	TGGACTCAAT	GAAGTGCACA	AGGCTGTGGG	GCTCCTGCTG	GAAACTCTAG	GGCCCCCACC .	, 900
10	CACTGGCCTG	CACCTGCAGA	GGGGAATCTA	CCGTGAGATA	TTATTCCTGA	CAATGGCTGC	960
	TCTGGGCAAG	GACCACGTGG	ACATAGTGGC	CTTCGATAAG	AAGTACAAGT	CTGCCTTTAA	1020
	CAAGCTGGCC	AGCAGCATGG	GCAAGGAGGA	GCTGAGGCAC	cccccccc	AGATGCCCAC	1080
15	TCCCAAGGCC	ATTGACTGCC	GAAAATGTTT	TGGAGCACCT	CCAGAATGCT	AGAGACCTTA	1140
	AGCTTCCCTC	TCCAGCCTAG	GGTGGGGAAG	TGAGGAAGAA	GGGATTCTAG	AGTTAAACTG	1200
20	CTTCCCTGTT	GCCTTCATGG	AGTTGGGAAC	AGGCTGGGAA	GGATGCCCAG	TCAAAGGCTC	1260
	CAAGCGAGGA	CAACAGGAAG	AGGGATCCAC	TGTTACCAAA	AGTCCTGATT	CCCCCATCAC	1320
35	CAACCTACCC	AGTTTGTTCG	TGCTGATGTT	GGGGGAGATC	TGGGGGGAGT	TGGTACAGCT	1380
25	CTGTTCTTCC	CTTGTCCTAT	ACCGGGAACT	CCCTCCAGG	GTACCCAÇAG	ATCTGCATTG	1440
	CCCTGGTCAT	TTTAGAAGTT	TTTGTTTTAA	AAAACAACTG	GAAAGATGCA	GAGCTACTGA	1500
30	GCCTTTGCCC	TGAATGGGAG	GTAGGGATGT	CATTCTCCAC	CAATAATGGT	CCCTCTTCCC	1560
	TGACGTTGCT	GAAGGAGCCC	AAGGCTCTCC	ATGCCTTTCT	ACCTAAGTGT	TTGTATTTTA	1620
25	TTTTAAATTA	. TTTATTCTGG	AGCCACAGCC	CCCTTGCTTA	TGAGGTTCTT	ATGGAGAGTG	1680
35	AGAAAGGGAA	GGGAAATAGG	GCACCATGGT	CCGGTGGTTT	GTAGTTCCTT	CAAAGTCAGG	1740
	CACTGGGAGC	TAGAGGAGTC	TCAAGCTCCC	CTTAGGAAGA	ACTGGTGCCC	CCTCCAGTCC	1800
40	TAATTTTTCT	TGCCTGCCCC	GCCTTGGGGA	ATGCCTCACC	CACCCAGGTC	CTGACCTGTG -	1860
	CAATAAGGAT	TGTTCCCTGC	GAAGTTTTGT	TGGATGTAAA	TATAGTAAAA	GCTGCTTCTG	1920
	TCTTTTTCAA	AANAAAAAA	AAAAAAAAACT	,			1950
45							

(2) INFORMATION FOR SEQ ID NO: 132:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 990 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 132:

TOGAAGATTT AAAATAGGTT TCATATTTCT CTTGAATATG AATATATAAG CTTGAATAAG

480

384

	CTTGAGTCCT TATTATTATG ARATTITCCT TATTATTTCT ACCARTGCTT CTTATATTAA	120
	AGCCTGATCT TTTTCATATT AGTATATGTA CATTAGCTGC CTGTGGATTA ACATTTCCAT	180
5	GARANGIANI TITGCANIGI TIGATOTTAA ACTITITIGIG TOTTTATATA AGGTATGCTY	240
	CTTTTAAGCA TGATATTTTT AACCAGAATA GTTGAAAGAC AATCTYCACC TTTTACTTGT	300
10	ATAITTACAT GTAATGIAAT TITTGATGCA TATTACGTCT TATTATTTAA CCAACCTATT	360
10	TTATTTTATC TAGGGCATTT TTCAGAAAGC CTTATTTTCT TGTATTAATC AAATATTTTT	420
	AYCAPTGTAT TTTCCYCTAT TAGTTAGKAA TACGKTACYC YAAATATATA TTGTGGSTAT	480
15	TTTCAGAATT GCAATATGCC TCCTTAATTT ATTAGAGGCT AACCTAAATT ATTACTTTTA	540
	CCACTTACTT GAAAATTCTG GAACTTTAGA ACATTTATTG TTTTATGCAT TTTAATTCTA	600
20.	CTTGTATTTT TACTACTCCT AAACATTATT ATTGTTTTAG ACAAGCCAAA ATATATNITG	660
20	TTATTATCTT ATYCTCCATT TCTTTCTGTA TTTTTATGCC ACTATGTATG CTCAATTTCC	720
•	TICHATGIGA TGAACCTAAT TCAGTACTIT TGTTTTTTAA TCTGTGCAGG TAGCCTGGCC	780
25	ATTAAATTIT TATTITIGGT TIGCTGAAAA AATTGTGTTT ATTTCTATAT GCATACTTAT	840
	GCATATAGAA TNCTAGGTNG ACATATTTTT AGTATTTATA AATGTAAAGT CATTWATTKG	900
30	GCTTCTATCA TYTCKGTKGA GAAATCAATT GTCAGCCCAA TAGTTTTTCA TYTTAAATTA	960
30	CNGAATTITT TCATGTTTCT GGTTTTAGGA	990
35	(2) THOROTTON TO TO ID NO. 122.	
	(2) INFORMATION FOR SEQ ID NO: 133:	
	(i) SEQUENCE CHAFACTERISTICS:  (A) LENGTH: 1720 base pairs	
40	(3) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLCGY: linear	-
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 133:	
	GTCTGATAAG CGACTGTGGT TATTCCCCTA AAGTTTACTT CAGCACTAAC ACTAGTGCTT	60
	CCGCTGGAGT TTGCAGTTTT CCAGCTTTAT ACAGGATTTT CCTTTGACTG GAAGAGTCAA	120
50	GGATATAGAG ACTCAALAGT GACATTTATT GTACAACATC AAGGGGAATA GGATACTCAT	180
	CALACTIGGGA TTATTCTTAT CAAAACATGG TCTTCTTTGA ATAAGAAAAA TACATAGTTG	240
55	GTTATTATGG ACTTAAAACT GTGTTAAATG GATATTCTGA TAAAATATTT GCTGCTCTGT	300
رر	AGASTGTGGA AAATCTGAGA ATATTAGCTT TACTCATCTT GAGCTTTGAG GATGTTCTCT	360

GTACGCCGAT GGTTTCATAT TAACTAAAAA AGCTGGGTAT TGTAAAATCT CATTTATAAA

AACTCAGATG AGAAGAAAAT TITCTTTGAT GGTGAGACTG TTGTCTTAGT TCAGGAAATT

120

	ATTTAATAAT CCTTTGTTAC CTGTGAATGA AGGAACTTTG TAATTCTGAT TTATCGTAAA	540
5	ACATGAGCCT TTCCAGAGTC AGCTTAGACA CTGTTGTCGC AAATAGCCAT GCTTTGCCTT	600
5	ATGCCAAGGA GGCCCAGAGG GAGGGCCTAG TCTTCCTCTG TTGCTGTACA TATATTGAAA	660
	TGCTTTTTT TTTTATTTTG CATTIGTTAT CTATAATGAG CTTTCTGAGC CCTGATATTA	720
10	TGTGAGACAA ACAGGAGITA TTGATGTTAT ACACTCCCTT CCATTCAGGA TTTTCTGCTT	780
	GGAGGGAAAT ATGTTGACCT TAGAGAATTG TGAATATTGT TGCAATTCTT GAATATATTA	840
15	CCATGIGAAT AATAGAGACT GTGTTGCTCT CTAGTATAAG CTATATTTAT TTTTGATTCA	900
15	TTTGAATTAC TAGTTATAAC TGGAGAAATT TTGTTACCTC TATCCTGGCT TGCCTGACTG	960
	GCTGTATAAT AGCAGCAGCC TCTTTTAGAG CATCTTAATG AAAACATGGA TGAAAGGAAT	1020
20	TAATGATGAT ATCTGCAGAC TGCGTAGAAA ATGGCTTTTG TTCCCAGCGT TAACATTTTC	1080
	TTCTCAATCA CATTTCAATG TTTGTGGAGA GTGGCAGATT CACACCAGAA ACACTAGGTG	1140
25	TTCATATCCA TAGCATGGAT GCAGAATAAG CAGTTGGGAG AGAAGCTTCT TCCTACCTGG	1200
25	TACTCCTCCC ATTCACCTCA GCCCAGCCCC AGACAGGCGT TAGCATTCAG TGTGGGCCCT	1260
	CAGGCAGCCC TGAAGCCTGG CTGGGTCATC AGATGGGGGC AGCCTGTGAC GGGCACCAGC	1320
30	GGCCTGATTC CAGGGAAGAG TTCCTGGAGG GTGTTGGCTG TTTTTGTTAG CTCAGTTTTT	1380
	TTCTGGGCTC CACCATTCCT AACTCCAGGT AGACAAGATA GATGTCACAC ACAACAATTT	1440
35	TAAAGTATTT TGCTTAGTGC ATTTTGTTTA TGATTGCAGT GTTTGTTTCT TATTTAATAG	1500
	GCTTTTTACT TCATTCTATT AAATTTTAGT GTTTAGAAGA GGCGGGTACT GTCACTGTGT	1560
	AAAATATGTA ATATTTTATA TGTTATACCA TGTCATATAT ACTTGCAATA TCAGACCTTG	1620
40	CATTCAATAT ACAATGCAAT TGACTCTTTG CAGACCTGCA TTTTTCAGTG AACAATAAAA	1680
	AGATTGTCTG GCACTCCAAA AAAAAAAAAA AAAAAAAAAA	1720
45		٠.
15	(2) INFORMATION FOR SEQ ID NO: 134:	
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 705 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 134:	
JJ		
	GGCACGAGGC CATCTGGGCT CATTCAGCAG GAAATAATGG AAAAAGCTGC AATATCCAGG	60

TGTTTACTAC AATCTGGAGG CAAGATCTTT CCTCAGTATG TGCTGATGTT TGGGTTGCTT

PCT/US98/11422

	•	
	GTGGAATCAC AGACACTCCT AGAGGAGAAT GCTGTTCAAG GAACAGAACG TACTCTTGGA	180
	TTAAATATAG CACCTTTTAT TAACCAGTTT CAGGTACCTA TACGTGTATT TITGGACCTA	240
5	TCCTCATTGC CCTGTATACC TTTAAGCAAG CCAGTGGAAC TCTTAAGACT AGATTTAATG	300
	ACTCCGTATT TGAACACCTC TAACAGAGAA GTAAAGGTAT ACGTTTGTNA AATCTGGGAA	360
	GACTIGACTG CTATTCCATT TTGGGTATCA TATGTACCTT GATGAAGANG ATTAGGTTGG	420
10	GATACTTCAA GTGAAGCCTC CCACTGGAAA CAAGCTGCAG TTGTTTTAGA TAATCCCATC	480
	CAGGITGAAA TGGGAGAGGA ACTIGTACTC AGCATTCAGC ATCACAAAAG CAATGTCAGC	540
15	ATCACAGTAA AGCAATGAAG AGCAGTTTTC CAATGAAAAC TGTGTAAATA GAGCATCAAC	600
	AAGTACAAAA TTCTTGTCTT AATTAGTGGG GGTATATAAA AATTCCTTGT AATGGTCAAA	660
20	TATTTTTAA AATTGACATT AATAAAGCAT ATTTTAAAAG TTTCT	705
20		
25	(2) INFORMATION FOR SEQ ID NO: 135:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 323 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
30	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 135:	
35	AGCACACAC TCCTTTAGTT GCTCCTAAGG TCATGTTCAA CATTCGTGGA GTGCATTTTC	60
<i>)</i>	TECTCAGGGA GCTTTCCCAG ACCCGGAATG TTTGGTGCTC ACAGACYCTG GCAAGGATCG	120
	GTATTGCTGT TCCTCAGTTT TGCCTGGGGA AATGGAGGST CAGTGACGTT CAGTGACGTG	180
40	CCCAGAGTCA TGCCATTGGC GGGTGGCCCA GKGMTCCAGG TCTCCAGCAC CCCTCGGCCC	240
	CCTCCTCACC AGGTCACATC ATCTCCTGGA TTAGAATCTG CTCACATAGT CTGTCCTGAA	300
45	ACGAAAAAA AAAAAAAAAA AAC	323
45		
50	(2) INFORMATION FOR SEQ ID NO: 136:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 582 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 136:	
	GGACGGAATG GTGCAACCCT CCTWAMTTTT CTKGKGCTGT TGACAACAGA GGGAGGGAGG	60
60		

	GAAAACATTT	TTYGTGGGAG	AATCCTACYT	CTGCAGSGGA	GCCCTTAAGC	GATKGATTTT	120
	GAATCTKGAC	CCTTTACCAA	CTAATTTTGA	AGGAAGATAC	CTTGGAAATA	TTTGGCATTC .	180
5	AGTGGGTTAC	TGAAACAGCA	TTAGTGAATT	CATCTAGAGA	ACTCTTTCAT	TTATTCAGGC	240
	AACAACTGTA	CAACTTGGAA	ACCTTGTTAC	AGTCCAGTTG	TGATTTTGGG	AARGTATCAA	300
0	CTCTACACTG	CAAAGCAGAC	AATATTAGGC	AGCAGTGTGT	ACTATTTCTC	CATTATGTTA .	360
·	AAGTTTTCAT	CTTCAGGTAT	CTGAAAGTAC	AGAATGCTGA	GAGTCATGTT	CCTGTCCATC	420
	CTTATGAGGC	TTTGGAGGCT	CAGCTTCCCT	CAGTGTTGAT	TGATGAGCTT	CATGGATTAC	480
l5 ·	TCTTGTATAT	TGGACACCTA	TCTGAACTTC	CCAGTGTTAA	TATAGGAGCA	TTTGTAAATC	540
	AAAACCAGAT	TAAGGTTTGA	CTGGTTTCAT	TTGATTTTTA	AG .		582

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#### (2) INFORMATION FOR SEQ ID NO: 137:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1021 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 137:

TTCGGCAGAG CCCTTGCGCG CTCTTGAATA CCTGCKTTCT GTAGCGCTAG TTCTCTTCAA 60 GATTTGCTTA GTGTCATTTC ATTTCGGTTT CTTTTCTCGC CATGTTTTTC TGTCGGAATT 120 ACGGTTCGTT TTGGTTCTAT GTACTCTCTA AAATGTTATC GTTTTTCATT TGTCTACTAA 180 TTTTCGTGCA TTTGTTACTA CTGAGTTTCT TAATATCTGA CTGGCCTCCG CCCACGGGCT 240 CTGCAGANCA TAAAATACTC AGGCTGATGG TAGTGCAGAG ACTCTCCCTC CTTGATCAGC 300 GCAAACGTTG GTCTGAGGCT TGAGGGATGG AGCAACATTT TCTTGGCTGT GTGAAGCGGG 360 CTTGGGATTC CGCAGAGGTG GCGCCAGAGC CCCAGCCTCC ACCTATTGTG AGTTCAGAAG 420 ATCGTGGGCC GTGGCCTCTT CCTTTGTATC CAGTACTAGG AGAGTACTCA CTGGACAGCT 480 GTGATTTGGG ACTGCTTTCC AGCCCTTGCT GGCGGCTGCC CGGAGTCTAC TGGCAAAACG 540 GACTUTUTCU TEGRASTICCAG AGCACUTTEG AACCAAGTAC AGCGAAGCCC ACTGAGTTCA 50 600 GTTGGCCGGG GACACAGAAG CAGCAAGARG CACCCGTAGA AKARGTGGGG CAGGCAGARG 660 AACCCGACAG ACTCAGGCTC CRGCAGCTTC CCTGGAGCAG TCCTCTCCAT CCYTGGGACA 55 GACAGCAGGA CACCGAGGTC TGTGACAGCG GGTGCCTTTT GGAACGCCGC CATCCTCCTG 780 CCCTCCAGCC GTGGCGCCAC CTCCCGGGTT TCTCAGACTG CCTGGAGTGG ATTCTTCGCG 840 TTGGTTTTGC CGCGTTCTCT GTACTCTGGG CGTGCTGTTC ACGGATCTGT GGAGCTAAGC 60 900

60

1020

1080

1140

1200

	AGCCTTAGAT AGCAGCAGAA GGCTTTTTGG ATTCTCCTCC TTGAAAAGAT TCTCAGTTAC	960
5	CAAACGTCTC CACCTAGAAA ATAAAAATAC ATTAAGATGT TGANAAAAAA AAANAAAAAA	1020
J	A	1021
		-
10	(2) INFORMATION FOR SEQ ID NO: 138:	
15	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 1777 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: double</li> <li>(D) TOPOLOGY: linear</li> </ul>	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 138:	•
20	CGGAAGATGA TGGCTTCAAC AGATCCATTC ATGAAGTGAT ACTAAAAAAT ATTACTTGGT	60
	ATTCAGAACG AGTTTTAACT GAAATCTCCT TGGGGAGTCT CCTGATCCTG GTGGTAATAA	120
25	GAACCATTCA ATACAACATG ACTAGGACAC GAGACAAGTA CCTTCACACA AATTGTTTGG	180
	CAGCTTTAGC AAATATGTCG GCACAGTTTC GTTCTCCCA TCAGTATGCT GCCCAGAGGA	240
20	TCATCAGTTT ATTTTCTTTG CTGTCTAAAA AACACAACAA AGTTCTGGAA CAAGCCACAC	300
30	AGTCCTTGAG AGGTTCGCTG AGTTCTAATG ATGTTCCTCT ACCAGATTAT GCACAAGACC	360
	TARATGICAT TGAAGAAGTG ATTCGAATGA TGTTAGAGAT CATCAACTCC TGCCTGACAA	420
35	ATTCCCTTCA CCACAACCCA AACTTGGTAT ACGCCCTGCT TTACAAACGC GATCTCTTTG	480
	AACAATTICG AACTCATCCT TCATTTCAGG ATATAATGCA AAATATTGAT CTGGTGATCT	540
	CCTTCTTTAG CTCAAGGTTG CTGCAAGCTG GGAGCTGAGC TGTCAGTGGA ACGGGTCCTG	600
40	GAAATCATTA AGCAAGGCGT CGTTGCGCTG CCCAAAGACA GACTGAAGAA ATTTCCAGAA	660
	TTGAAATTCA AATATGTGGA AGAGGAGCAG CCCGAGGAGT TTTTTATCCC CTATGTCTGG	720
45	TCTCTTGTCT ACAACTCAGC AGTCGGCCTG TACTGGAATC CACAGGACAT CCAGCTGTTC	780
	ACCATGGATT CCGACTGAGG GCAGGATGCT CTCCCACCCG GACCCCTCCA GCCAAGCAGC	840
<b>50</b>	CCTTCAAGTT CTTTTATTTC TGGGTAACAG AAGTAGACAG ACAGGTTACT TGGTGTATCT	900
50	TCTGTTAAAG AGGATTGCAC GAGTGTGTTT TCCTCACACA CTTTGATTTG GAGAATTGGT	960

GCTAGTTGGC AATAGATAAC TCAGCGTAGA TAGTATTGCA AAAAGGGGAG GAAATACACA

ACAATAATAA ATGTAAAAAC CTGCTATTCA ACATGCAGTT TTATTTCGAR GCCAAAAATC

TAGAGCTTTC CCAAGATCCT GTTGCCTTAG GCACATNCAC ACTTCAACAG TGCACACTAT

CCAACAGTGC ACACTATTCA ACAGTGCACA CTATTCAAAA GCGTAGACTA TTTTTTTGCA

	TGTTCAAGAT ATTTGTTTTG GTCTTATGTG TGTGTGAGAG AGAGAGATTC CTTTGACATT	1260
	AAGGAGCATC AATGAGAAAA GATGATGAGG CAGGAATTAA TAAAGAAATG AAGTCGTGTG	1320
5	TGTTTGGTTG CCTGTCAGAG GGCACACAAT TTCATAAACA CCATGCCTGG ACAATTTGAT	1380
	ATTAATATTT AACACCTCTG CATCTTTTTC TTAAAAAAGA ATATGGGCCA GATACAGTGG	1440
10	CTCACATTTG TAATCCCAGC ACTTTGGGGA GCCAAGTTAG CAGAATCCCT TGAGCACAGG	1500
10	AATCTGAAAC CAGCTTGGGC AACATAGTGA GATCCCATCT NTACAAAAAA CTTAAAAATT	1560
	AGCCAGGCAT GATGGCACAT TCCTGTAGTC CTAGCTACTC AGGAGGCTAA GGTAGGAGGA	1620
15	TTGCCTGAGC CCAGGAGTTC AAGGCTGCAG TGAGCTAAGN ACGTGCCAGT ACACTCCAGC	1680
	CTGAGCCACA AAGTGAGACC CTGTCTCGCA AAAAAAAAAN TTAAAAAGTC GGGGGGGGC	1740
20	CCGGTACCCA AATCGCCGGA TATGATCGTA AACAATC	1777
20		
	(2) INFORMATION FOR SEQ ID NO: 139:	
25		
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 643 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
30	(D) TOPOLOGY: linear	
•	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 139:	
35	TITTTTTTT TITTTTTTT TITTTTTTT TITTTTTTGGG AATGAGAAAA TAACTTTATT	60
33	TTCATTGTGG GGAGCGGGCC GATGTCCAGC CTCAGAACTT CTGGAACTGC TTCTTGGTGC	120
	CGGCAGCCTT GGTGACCTTG AGCACGTTGA AGCGCACTGT CTTGCTCAGA GGCCGGCACT	180
40	CGCCCACTGT GACGATGTCA CCGATCTGGA CGTCCCTGAA GCAGGGGGAC AGGTGTACAG	240
	ACATGITCTT GTGGCGCTTC TCGAAGCGGT TGTACTTGCG GATGTAGTGC AGATAGTCTC	300
15	GGCGGATGAC AATGGTCCTC TGCATCTTCA TCTTGGGTCA CCACGCCAGA GAGGATCCGC	360
45	CCTCGAATGG ACACATTACC AGTGAAGGGG CATTTCTTGT CAATGTAGGT GCCCCTCAAT	420
	AGCCTCCTTG GGGTGTCTTT GAAGCCCAGA CCGATGTTCT TGTTAGTAAC CCGCGGGAGC	480
50	TTCTCCTTGC CAGTTTCTCC CAGCAGGACC CTCTTCTTGT TTTGAAAGAT GGTCGGCTGC	540
	TTTTGGTAGG CACGCTCAGT CTGAATGTCC GCCATCTTCT CGTGCCGMAY TCCTGCAGCC	600
55	CGGGGGATCC ACTAGTTCTA GAGCGGCCGC ACCGCGGTGG AGC	643
JJ	•	

<sup>(2)</sup> INFORMATION FOR SEQ ID NO: 140:

1:1	CECTEMOE	CHARACTERISTICS
(1)	SECUENCE	CHARACIERISIICS

(A) LENGTH: 1220 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 140:

10	GGCACGAGGA	TGATAGACCT	ACTGGAGGAA	TACATGGTTT	ACAGGAAGCA	TACCTACATR	60
10	AGGCTTGATG	GCTCATCCAA	GATCTCGGAG	AGGCGAGACA	TGGTTGCTGA	TTTTCAGAAC	120
	AGGAATGACA	TCTTTGTGTT	CCTGTTAAGC	ACACGAGCTG	GAGGACTGGG	TATCAATCTC	180
15	ACTGCTGMAG	ACACAGTGCA	TTTTCTATGA	TAGCGACTGG	AACCCCACTG	TGGACCAGCA	240
	GGCCATGGAC	AGGGCCCACC	GCTTAGGGCA	GACAAAGCAG	GTTACTGTGT	ACCGGCTCAT	300
20	CTGTAAAGGC	ACCATTGAAG	AACGCATTCT	GCAAAGAGCC	AAGGAGAAGA	GTGAGATTCA	360
20	GCGGATGGTG	ATTTCAGGTG	GGAACTTCAA	ACCAGATACC	TTGAAACCCA	AAGAGGTGGT	420
	TAGTCTTCTT	CTAGACGACG	AAGAGTTGGA	GAAGAAACGT	ATGTACTCTA	AACCTCTATA	480
25	CACTCCCCTC	ACGTATCTGA	GAATGGAAGA	GGTACTTGGS	TGTGTGCCAA	GGGTTAGGCA	540
	AAGCCAGAGG	CTGTATTTAG	GGAAAGTATT	TTTGTGCTCA	TATTTTATAT	AAAAACCCAA	600
30	ACAAGAATGT	GTTTGTAGGC	CAGGCGTGGT	CCCTCCCCCC	TCTAGTCTCA	CCATTTCGGG	660
50	ARGCCAAAGT	GGGCAGATCA	CCTGARGTCA	GGARTTTGAG	TTTGARACCA	GCCTGGCCMA	720
	CGTTGTGAAA	CCCCACCTCT	ACTARGARTA	CSGAAAATTG	GTTGGGCATG	GTGGCGGGCA	780
35	CCTGTAATTC	CAGCACTTTG	GGAGGCTGGG	GCAGAANAAT	TGCTTGAGCC	CAGGAGGTGG	840
	AGATTGCGGT	GAGCCGAGAT	YGTGCCATTG	CAMTCCAGCC	SGGGCAATAA	GAGTGAAAYT	900
40	CCATCTTTTA	AAAACAAACA	ААААСААААА	ACACAAGACG	GCTCACACCT	GTAATCCCAG	960
40	CACTTTGGGA	RGCCGARGCA	GGTGGATCAC	GARGTCAGGA	GTTCCAAGAC	TAGCCTGGCC	1020
	AACCTGGTGA	AGCCCCGTCT	СТАСТАААА	TACMAATATT	AGTCGGGCGT	GGTGGTGGGC	1080
45	ACGTGTAATC	CCAGCTACTC	GGGAGGCTGA	GGCAGGAGAA	TCCCTTGAAG	CTAGGAGGCA	1140
	GAGGTTGCAG	TGAGCCAGGA	TCGTGCCATT	GCACTCCAGC	CTGGACAACA	AGAGCAAGAT	1200
50	TCCATCTCAA	AAAAAAAAA					1220
~							

(2) INFORMATION FOR SEQ ID NO: 141:

55

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 721 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(XI) SEQUENCE DESCRIPTION SEQ IS NO. 111.	
5	AATTCGGCAC GAGCCAGGTT AGCCGGAAGG GCAGCTCTCC AGGCCCTGCC CACCCCACAG	60
5	GGGGCTCCTT ATGCACAGCG GGGCGTCTCC TTGTGGCCAT AGAAACGGAA CTGGCTCTTT	120
	TCAACAGTGC TGCAAGAGGA TGGTTATTTA ACGCTGGCCC CCAAGGAGGA AAGGCACAGA	180
10	CYTTCCTCCC TCCTGGAACA TCCAAGGGCA CTGGATCCTC TGTGTCCCTC TGAGATGGGG	240
	TGCCACTCCA GCAAGAGCAC CACGGTGGCA GCTGAGTCCC AGAAGCTTGA AGAAGAGYGC	300
15	GAGGGAAGAG AGCCAGGTCT GGAGACCGGC ACCCAGGCAG CAGACTGCAA GGATGCCCCG	360
13	CTGAAGGATG GAACCCCTGA GCCAAAGAGC TGAAATGCCT CTCTCCAGAG TCGGACCCTC	420
	ACCTCYTTCC TGGAACTGCC TTTGGCCCCA GAACCATGAG ACAATCCCCA CCCTGAGAAG	480
20	CTCCGATCAC TGGGAGGAGA GAGAAAGCCT CCAGCTTTGG GATTCAGGCT TCAGAAGTTT	540
	TTAGCAGCCT TTGCTCATTG GAGAGGTGGG GAAAGGATAA AGTTCTTATA AGGAAATCCC	600
25	TAATTTCCCC CAGCTCCTCC CCNCCNGAAG AAGGAACNAA AGAAAGTTCC TTCCACACGT	660
23	TTTGTTGGAA ACTTTTCCCT TGCCAACTTT CCTTGGATTG CCAGAACAAA GCCCTCCAGA	720
•	A	721
30		
	(2) INFORMATION FOR SEQ ID NO: 142:	
35	(i) SEQUENCE CHARACTERISTICS:	1
	(A) LENGTH: 1468 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 142:	
	ATGAATTAAT GTTTATAAAT GACTGTACTG AATTTAAAAC CGTACAGTTT CATTTGCATT	60
45	TTGACATTAC TITATTATAC ATTITGCATT TAAAAGGCTG CACCAGTTGG CTTTTCTTCT	120
	GTTTTATTCT CAAAATATAG AGATTCTGTG ATTTATTTGC CCTGTTTATG GATTAAAAAG	180
	AAAATTCTAA TATAAAGCAT TTCAATAGGA TGCATAGGTA TATTACGTTT TTTAAATGCT	24
50	TTAGATCTGT GATTCTTGAC TTACTATTTA TTTTATCCCC TTTAAGTCAG GGATGCTTTA	30
	TICTATITTA AAGCACTTAT GAGTTACATG TIGTAATCAA GITIGCACAA TATATITATC	36
55	TATATGAGGA ACCCATAAAT GAATAGCTAA TTTTTAAAAT GCCATTAAAA TGCATGAAAT	42
	KCTTATTAAA ACCTTACTAT ACTATTTCTT CAAGGCAAGT AAATTGACCA TGRGRAAAGR	48
	ACACAGTTAT TAAACACTGT TGACAGGAAA ATTCTCCTTG ATAACATAGG ACAATTAATG	54
60		

	GAAAAAAAA TTCTCATTAT TTGCAAAGAA TGAACAAGTT AATGAACAAA CAAACTAGAT	600
	TTGGTATGTT TTCAGCTTTT GTATCATGTT TAATTGTTTA ATTTGGTTGA AAAACTGCAG	660
5	TTGAGAAATC AGATAGCAAT ATAGACATTC ACAGCAGCTC TGTGGATACC ATGTAATTGT	720
	CAGGTAATTT CAGAATGTTG AAAATTATTC AGTGCAGCCC TCATAGTATC ATACTTGAAG	780
	AAATTGATTA CAGTTCCACT AAATTGTTGA AGATAAATTA TTTTTAAAGG TTATGAAAAC	840
10	TAAGTTATAT TAATTCATAT GTTTGATTTT TAAATCCCAC CTCCTCAAGC TATCCAATTT	900
	NCTGACTTTG AAAATAACCA TGAGAGATGC CACATTTCTC TCTGGGAAAC TACCACTCAA	960
15	AGAATAATTG TTAAAAATTA AGCTTTTAGG TATTAGAAGC TGTTATAAAG TATAAAATTA	1020
	AGATATAAGC AGATCACATG TAAATCATTC CTAAAGCACA AGAAAAGAAT GTGCCTTGAT	1080
20	GTACATATAT TACTAAGTTG CCTCTCCCAG TTTACTTTAA AAATGGCTTT AAGGATAAAG	1140
20	AATAAATGTG ATAGCTGTGC ATGCATTATA TATTTGCATT TGCAAATTTC CCATTGTTTT	1200
-	AACAGCTGTG TGGCTGACTT TCAATTTTAA GACGTGAATT GACATACAGC CCATAACTTT	1260
25	ATAATGGCTG CTCATTTATC TTATCTTTCA GTTAGTGGAA AAACATTTCA ACCTGACTAA	1320
	AATTIGGAAT TGTGTCTTTT ATGTTCCATC CTCTGTTGTT ACTAGATTTA GTTTAAAAAT	1380
30	TGTGTATGAC CATTAATGTA TGTCATAAAC ATGTAAATAA AAGATGTTGA ATCTTGTTGA	1440
30	AAAGCAWRAA AAAAAAAAA AAACTCGA	1468
35	(2) THEODIS STOLE SEC. TO NO. 147.	
	(2) INFORMATION FOR SEQ ID NO: 143:	
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 300 base pairs	
40	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 143:	60
	TGAATTITTT GCCAAACTTA GTAACTCTGT TAAATATTTG GAGGATTTAA AGAACATCCC	
50	AGTTTGAATT CATTTCAAAC TTTTTAAATT TTTTTGTACT ATGTTTGGTT TTATTTTCCT	
50	TCTGTTAATC TTTTGTATTC RCTTATGCTC TCGTACATTG AGTACTTTTA TTCCAAAACT	
	AGTGGGTTTT CTCTACTGGA AATTTTCAAT AAACCTGTCA TTATTGCTTA CTTTGATTAA	•
55	AAAAAAAAA AAAAAAAAAA AAACCCCNAG GGGGGGCCG GGTNCCCAAT CCCCCCCAAA	300

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2243 base pairs(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 144:

10	mecemeceena.	CCTGCAGATT	CMCC A C A C M A		CITCE A C CCTIC	CAMMCCMMCM	. 60
	TCCCCTTCCT	AGCTCCATGG	GACTCGCCCC	AAGACTGTGG	CTTCAAGGAC	CACCAGCCCC	120
15	TTACTCTTCA	AGCCCTGACT	GTGGAGTTGG	TAGATGCCTC	TGATCCTCAG	TATTCTCTCT	180
	GGCAATGTTC	CACGGCTTCT	CCTTCCTGGG	AGCTGGCTCC	ATAACTTGAT	TTTCCCCAAA	240
	CGTGTTGCAA	TCCCTGCTGC	CCCTTAGCCA	CCCAGGGTCT	TGTGTGGGTA	TGAGTGTAGA	300
20	GGATGGGGGT	ATGCCAGGCC	TGGGCCGTCC	CAGGCAGGCC	CGCTGGACCC	TGATGCTACT	360
	CCTATCCACT	GCCATGTACG	GTGCCCATGC	CCCATTGCTG	GCACTGTGCC	ATGTGGACGG	420
	CCGAGTGCCC	TTYCGGCCCT	CCTCAGCCGT	GCTGCTGACT	GAGCTGACCA	AGCTACTGTT	480
25	ATGCGCCTTC	TCCCTTCTGG	TAGGCTGGCA	AGCATGGCCC	CAGGGGCCCC	CACCCTGGCG	540
	CCAGGCTGCT	CCCTTCGCAC	TATCAGCCCT	GCTCTATGGC	GCTAACAACA	ACCTGGTGAT	600
30	CTATCTTCAG	CGTTACATGG	ACCCCAGCAC	CTACCAGGTG	CTGAGTAATC	TCAAGATTGG	660
30	AAGCACAGCT	GTGCTCTACT	GCCTCTGCCT	CCGGCACCGC	CTCTCTGTGC	GTCAGGGGTT	720
	AGCGCTGCTG	CTGCTGATGG	CTGCGGGAGC	CTGCTATGCA	GCAGGGGGCC	TTCAAGTTCC	780
35	CGGGAACACC	CTTCCCAGTC	CCCCTCCAGC	AGCTGCTGCC	AGCCCCATGC	CCCTGCATAT	840
	CACTCCGCTA	GCCTGCTGC	TCCTCATTCT	GTACTGCCTC	ATCTCAGGCT	TGTCGTCAGT	900
40	GTACACAGAG	CTGCTCATGA	AGCGACAGNG	GCTGCCCCTG	GCACTTCAGA	ACCTCTTCCT	960
40	CTACACTTTT	GGTGTGCTTC	TGAATCTAGG	TCTGCATGCT	GCCGCCGCT	CTGGCCCAGG	1020
45	SCTCCTGGAA	GGTTTCTCAG	GATGGGCAGC	ACTCGTGGTG	CTGAGCCAGG	CACTAAATGG	1080
	ACTGCTCATG	TCTGCTGTCA	TGAAGCATGG	CAGCAGCATC	ACACGCCTCT	TTGTGGTGTC	1140
	CTGCTCGCTG	GTGGTCAACG	CCGTGCTCTC	AGCAGTCCTG	CTACGGCTGC	AGCTCACAGC	1200
50	CGCCTTCTTC	CTGGCCACAT	TGCTCATTGG	CCTGGCCATG	CGCCTGTACT	ATGGCAGCCG	1260
	CTAGTCCCTG	ACAACTTCCA	CCCTGATTCC	GGACCCTGTA	GATTGGGCGC	CACCACCAGA	1320
						GCCTTGTGAG	
	AAAAGCTGGA	GAAGTGAGGG	CAGCCAGGTT	ATTCTCTGGA	GGTTGGTGGA	TGAAGGGGTA	1440
	CCCCTAGGAG	ATGTGAAGTG	TGGGTTTGGT	TAAGGAAATG	CTTACCATCC	CCCACCCCCA	> 1500
						GAGAAATAAC	
60	racento 11CT	ICCHONCIAM	UGULI INUGG	IMMITTAL	.202.00001	- man a marking	1300

	CCCATCCTTG	TTGGGCAGCT	CCCTGCTTTG	TCCTGCATGA	ACAGAGTTGA	TGAAAGTGGG	1620
	GTGTGGGCAA	CAAGTGGCTT	TCCTTGCCTA	CTTTAGTCAC	CCAGCAGAGC	CACTGGAGCT	1680
5	GGCTAGTCCA	GCCCAGCCAT	GGTGCATGAC	TCTTCCATAA	GGGATCCTCA	CCCTTCCACT	1740
	TTCATGCAAG	AAGGCCCAGT	TGCCACAGAT	TATACAACCA	TTACCCAAAC	CACTCTGACA	1800
10	GTCTCCTCCA	GTTCCAGCAA	TGCCTAGAGA	CATGCTCCCT	GCCCTCTCCA	CAGTGCTGCT	1860
	CCCCACACCT	AGCCTTTGTT	CTGGAAACCC	CAGAGAGGGC	TGGGCTTGAC	TCATCTCAGG	1920
	GAATGTAGCC	CCTGGGCCCT	GGCTTAAGCC	GACACTCCTG	ACCTCTCTGT	TCACCCTGAG	1980
15	GCTGTCTTG	AAGCCCGCTA	CCCACTCTGA	GGCTCCTAGG	AGGTACCATG	CTTCCCACTC	2040
	TGGGGCCTGC	CCCTGCCTAG	CAGTCTCCCA	GCTCCCAACA	GCCTGGGGAA	GCTCTGCACA	2100
20	GAGTGACCTG	AGACCAGGTA	CAGGAAACCT	GTAGCTCAAT	CAGTGTCTCT	WTAACTGCAT	2160
	AAGCAATAAG	ATCTTAATAA	AGTCTTCTAG	GCTGTAGGGT	GGTTCCTACA	ACCACAGCCA	2220
	ŸYYYYYYYY	AAAAAAACTC	GAG ·				2243

## (2) INFORMATION FOR SEQ ID NO: 145:

30

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1082 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

35

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## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 145:

GCCAAGCTCT	AATACGACTC	ACTATAGGGA	AAGCTGGTAC	GCCTGCAGKT	ACCGGTTCCG	60
GGAATTCCCG	GGTCGACCCA	CGCGTCCGCT	TCCGTGTGTC	AAAATCCTCA	CCTCCTTCAT	120
AACCATCTCC	CACAATTAAT	TCTTGACTAT	ATAAATTTAT	GGTTTGATAA	TATTATCAAT	180
TTGTAATCAA	TTGAGATTTC	TTTAGTGCTT	GCTTTTCTGT	GACTCAACTG	CCCAGACACC	240
. TCATTGTACT	TGAAAACTGG	AACANCTTGG	GAATGCCATG	GGGTTTGATA	ATCTGCCAGG	300
GACATGAAGA	GGCTCAGCTT	CCTGGGACCA	TGACTTTGGC	TCAGCTGATC	CTGNACATGG	360
GAGAACAACC	ACATTTTCT	TTGTGTGTGC	TTCTAGCAGC	TGTTCGGGAG	GACCKTGACC	420
CAAYAGTGTT	CCCATGCTGT	TTCTTGTGAA	ATGCTCTCGG	CTATGTAGCA	GCTTTTGATT	480
CCCTGCATAC	CCTAGGCTGC	TGCCCCTATC	CTGTCCCTTG	TTTATAACAT	TGAGAGGTTT	540
TCTAGGGCAC	ATACTGAGTG	AGAGCAGTGT	TGAGAAGTCG	GGGAAAATGG	TGACTACTTT	600
TAGAGCAAGG	CTGGGCATCA	GCACCTGTCC	AGCTCTACTT	GTGTGATGTT	TCAGGAACTC	660
AGCCCCTTTT	TCTGCCTAGG	ATAAGGAGCT	GAAAGATTAA	CTTGGATCTY	CTAATGGTCC	720

	AAATCTTTTG GTCACAATAA AGAGTCTCCA AATTAGAGAC TGCATGTTAG TTCTGGATGG	780
_	ATTTGGTGGC CTGACATGAT ACCCTGCCAG CTGTGAGGGG ACCCCGTTTT TAAGATGCAT	840
5	GGCCAAGCTC TCTGCAAATG GAAATGCTTA CACTGGGTGT TGGGGATGTT TGCTACCTCC	900
	TGCTATTTTT GTGGTTTTGG TTCTCCCACT ATGGTAGGAC CCCTGGCCAG CATTGTGGCT	960
10	TGTCATGTCA GCCCCATTGA CTACCTTCTC ATGCTCTGAG GTACTACTGC CTCTGCAGCA	1020
	CAAATTTCTA TITCTGTCAA TAAAAGGAGA TGAAAATAAA AAANAAAAA AAAAAACTCG	1080
	NG	1082
15		
20	(2) INFORMATION FOR SEQ ID NO: 146:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 4313 base pairs	
25	(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 146:	
30	CAAGCTGGTT TGAAACTAGG GGTCGGGCTC GGCCGTCGTC GTTGTTTGTC GCCGCATCCC	, 60 ,
	CGCTTCCGGG TTAGGCCGTT CCTGCCCGCC CCCTCCTCTC CTCCCTTCGG ACCCATAGAT	120
	CTCAGGCTCG GCTCCCCGCC CGCCGCAGCC CACTGTTGAC CCGGCCCGTA CTGCGGCCCC	180
35	GTGGCCACCA TGTCCCTGCA CGGCAAACGG AAGGAGATCT ACAAGTATGA AGCGCCCTGG	240
	ACAGTCTACG CGATGAACTG GAGTGTGCGG CCCGATAAGC GCTTTCGCTT GGCGCTGGGC	300
40	AGCTTCGTGG AGGAGTACAA CAACAAGGTT CAGCTTGTTG GTTTAGATGA GGAGAGTTCA	360
	GAGTTTATTT GCAGAAACAC CTTTGACCAC CCATACCCCA CCACAAAGCT CATGTGGATC	420
	CCTGACACAA AAGGCGTCTA TCCAGACCTA CTGGCAACAA GCGGTGACTA TCTCCGTGTG	480
45	TGGAGGGTTG GTGAAACAGA GACCAGGCTG GAGTGTTTGC TAAACAATAA TAAGAACTCT	540
	GATTTCTGTG CTCCCCTGAC CTCCTTTGAC TGGAATGAGG TGGATCCTTA TCTTTTAGGT	600
50	ACCTCAAGCA TTGATACGAC ATGCACCATC TGGGGGCTGG AGACAGGGCA GGTGTTAGGG	660
30	CGAGTGAATC TCGTGTCTGG CCACGTGAAG ACCCAGCTGA TCGCCCATGA CAAAGAGGTC	720
	TATGATATTG CATTTAGCCG GGCCGGGGGT GGCAGGGACA TGTTTGCCTC TGTGGGTGCT	780
55	GATGGCTCGG TGCGGATGTT TGACCTCCGC CATCTAGAAC ACAGCACCAT CATTTACGAA	840
	GACCCACAGC ATCACCCACT GCTTCGCCTC TGCTGGAACA AGCAGGACCC TAACTACCTG	900
	GCCACCATEG CCATGGATEG AATGGAGGTG GTGATTCTAG ATGTCCGGGT TCCTGCACAC	950

	CTGTSGCCAG GTTAAACAAC C	ATCGAGCAT	GTGTCAATGG	CATTGCTTGG	GCCCCACATT .	1020
	CATCCTGCCA CATCTGCACT G	CAGCGGATG	ACCACCAGGC	TCTCATCTGG	GACATCCAGC	1080
5	AAATGCCCCG AGCCATTGAG G	ACCCTATCC	TGGCCTACAC	AGCTGNAAGG	WGAGATCAAC	1140
	AATGTGCAGT GGGCATCAAC T	CAGCCCGAA	YTGTCGCCAT	CTGCTACAAC	AACTGCCTGG	1200
	AGATACTCAG AGTGTAGTGT T	GGTGGCGCT	GTGCCCACGA	GCAGGGCT	TTTGTATTTC	1260
10	CTGCCTCTGC CCCACCCCCA A	AGTAAGAAG	AAACATGTTT	CCAGTGGCCA	GTATGTCTTT	1320
	CATTGCTTTG CACCCACTGT T	ACCAGAAGC	TGCTCTAGGA	GTTCCTGGCC	AGTCACCCCA	1380
15	TCGCCCTCTG TGGCAGACTC A	GTGCTGTGT	GCCCCTCCT	CAGCCCAGGG	CTGAGTTTTA	1440
	AGATTTICTC TCCTTTCCTC T	TCTCCTTTG	GTTCCTCAAT	TAAAAAATGT	GTGTATATTT	1500
30	GTTTGTCAGG CGTTGTGTTG A	GGAGCAGTT	CACGCACTGG	CTGTGTCTAT	TCCTCTGCCC	1560
20	AGGTGTCTCT GTTTGCTGCC C	AAKGYWKKT	TTTCATGTCT	CGTCCATGTC	CATGTTCGTG	1620
	TTAGCACTWA CGTGGGAACA A	ATACCAATT	TGTCTTTTCT	CCTAGTATCA	GTGTGTTTAA	1680
25	CAAATTTTAA CTTTGTATAT T	TGTTATCTA	TCAGGCTAAT	TTTTTTATGA	AAAGAATTTT	1740
	ACTOTOCTGC TTCATTTCTT T	CTCTTATAG	TCCTCCCTCT	TTGCACCTTC	TTCTCTTCCC	1800
20	TCAGTGCCTG GAGCTGGTAC T	receccere	GCCCCATGAG	CAGTITGCCT	TCTTGAGTCA	1860
30	CTGCCTGTGT AGTACATACC T	rgaccgggag	TCCAAACCAC	CTTGGTGCTC	TGAAGTCCAC	1920
	TGACTCATCA CACCTTTCTT A	AGCCTGGCTC	CTCTCAAGGG	CATTCTGGGC	TTGTAAACAG	1980
35	ACATAGGAAG CCTCTGTTTA C	CCTGAAGCA	CCACTGTCCA	GCCCATTGGT	TCCCACTGGC	2040
	AGCATGGTAG AGCTGAGAGA	AACAGGCTCT	CAGGGTACCT	GACTTGAGGG	GAATCGTTTC	2100
40	ATGAAGCTGA ACTTCAAGCA	PATTTCCAGT	ACATTCTTTC	AGAGTCTGTT	TTTCCATCCA	2160
40	AATATAAGCC CCAGGCCATT (	CCACTTAGTG	TCTTTTCAAT	GATAGGCAAG	AATGATATCT	2220
	GAGTTGAACT TCGGTGCTTC 1	TGTTGTTTGA	GTTTACTGTG	CCTGGTGGTA	TATTGGGCAT	2280
45	TCTTTGGATT GAGTGTTCTG	AGÇTGAGAGA	GTCTTCCCGA	GCATCCTGT	CTGTGCTTCC	2340
	AACCCTGAAC AAGACCTTAC	atgagagatg	GACTGATGGA	CTGCGGCAAT	ccreecter	2400
50	CAAGTGGATA GATAGTTAAA	AAGCATTATA	CTGTGGGTAA	TGAAAAGGA	A GGAAAAAAA	2460
30	AGAAGGAAAA GGAATTATAG	ACCCCCAGGG	TCAGCCAGTT	AAGAGCTCT	A CCCACACCTG	2520
÷	TCAACCCCTC TCTCCCCCAG	TTTAGGTTCT	GAGCAGTATI	GGACTTGTA	CCTGCAGTTG	2580
<b>55</b> <sup>^</sup>	TCTTTTGACT TGCAGGCCGC	AGTGTCTTTC	TGTTATGTG	ATGAGTTCC	A TGGAGGGGCA	2640
	TATGTGTGAT TCCACCGTTA	GATGAGCCCT	TGGGGCAGG	AGTTTGGGA	r grgctcttgg	2700
60	GGGAAAGTTG GCTGTTTCCT	TGCGCTCTGC	TCCTACCCG/	A AGTTTTTAA	TCCCTCTGAA	2760

	TIGCTCATCT GAGATTAGTA GAGTAGCAGG CCTGAAGGAT GATGGTTTTG TCCTCTTTGG	2820
	TTCTCACCTG CTTGAGAAGT AAAACAGTAA CTTTGTTCTT CTGGGCCCTT AAGCTTTTTT	2880
5	GGTTAAGTCT TCCTTTTCAG AAGTAGATGT CATTATATGC CAAAAGTCTA GCTCTTTGCT	2940
•	TTACCATACA GGGACCTGTC CCAAAGAAAA AGGCTCTTTT TTTAGCCAGC ATATTTCCCC	3000
10	TTCTACCCTT TTACTTTGTT GTTCTGATTT TAGGACTCTG GCTGGCCATG TGCTTGTGGT	3060
10	TGCCTCTCCT GCATTTGCCA CTGGATTTGC ACTGCATCGT TTGGAGATAC AAAGCGAGCA	3120
	GTTCTTGGTC AGAACCCTCC TCTGCTTTTC ATTGTGTTTG ATAATGGTTA CTGGGTCCTT	3180
15	CTCTCAAGGG TAGCAAGGCC AAGCTGATGG CTGCTTGTTT AGGAGGCCAT CAGTTCCTTC	3240
•	CTGTGGAGAA GGGTCTGAAA TGGAAGTCAG TGGTAGAAGG GGCTGGTCTG CTGGGCAGGG	3300
20	CTTACATCCA CTGAGTTCTA AGATTCCTTT CCTGATCTGC ACCTACGCCT GGTCTGTATG	3360
20	GTGGAATTTG TCAGCTGGAA CTCAGAAACA ACAACTTGAA AAAAAAATAA TAATTAGAAC	3420
	ATATTTGCAT AAGATAGCTA TTTACTCTGG AAACCAACAA CTTTTGAGAT TTCCCTTGCC	3480
25	CTGTGGACGC CCAGCTCCTG TCATCCTTCC TTAGGTCCTG CAGTACAGTC TTCCCCTGAA	3540
	TGCCACCGGG GACCCAGGGG GACTCCACCC CCCTAAGCAA GCACACACAT ACTCACAGTT	3600
30	GATGAGTTGC TGGTCTTTGA GTCCCAGCTC TCTTACCCTC CCTTTACTCC ACCAGCCCGA	3660
20	CGACCCATGA CTGAGGAGGG GATTTCTACA GTCTCAGGAT TTAGAAAGTC TGTAAGCCAT	3720
	CCATGCTCCA GAAAGCACCG ATCTGTTGTA GTTGCAAAAA CAACTCTGTA ATTTGTTGAG	3780
<b>35</b> .	GTTCTCAAAC TGACAGCCAG CGAGACTGGG TGGGAGGCCC TGGATCTGTT CTCCCTGACT	3840
	GCGGGAGGAG CAGCCACTAG GACTTTAGCA GGAAGCCCAC ATGGAGGCTC CGCCAGGCTG	3900
40	TGGCCCAGCT GGTGATGGCC CTTTTGCTCC TGGCAGCCTG AGGCACAGCT GCCTGTATTG	3960
10	TCCTCATCTG TTCTGACTGA AGGATGGAGG TGCTGAATAA ATTAGGCCTC AGGCNTCTAC	4020
	CACCAGAGAG CTGGAGAATG GGTCCACGTC ATTCAAGGAC CTGAATTTTT TATGCTCAGG	4080
45	AGCATTGGAA TCCTCTTCTT CCAGGGAGGA ATTAGCCTGC AAGGTTAGGA CTTGAAGAGG	4140
	GAAGGTATTT AATAACTGGG CGAGGATGGG TGTGGTGGCT CACACCTGTA ATCCCAGCAT	4200
50	TTTGGGAGGC TGAGGTGGCC AGATCCCAAG GTCAGAAGAT CGAGACCATC CTGGCTAACA	4260
50	TGGTGAAACC CCATCTCTAC TAAAAATACA AAATTAAATT	4313

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1183 base pairs

(B) TYPE: nucleic acid

<sup>(2)</sup> INFORMATION FOR SEQ ID NO: 147:

(C)	STRANDEDNE	ESS:	double
(D)	TOPOLOGY:	line	ear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 147:

5							
5	GGCAGAGCCT	CAAGCTGACT	TGGATTATGT	GGTCCCTCAA	ATCTACCGAC	ACATGCAGGA	60
	GGAGTTCCGG	GCCCGTTAG	AGAGGACCAA	ATCTCAGGGT	CCCCTGACTG	TGGCTGCTTA	. 120
10	TCAKWYGGGG	AGTGTCTACT	CAGCTGCTAT	GGTCACAGCC	CTCACCCTGT	TGGCCTTCCC	180
	ACTTCTGCTG	TTGCATGCGG	AGCGCATCAG	CCTTGTGTTC	CIGCTICIGT	TTCTGCAGAG	240
15	CTTCCTTCTC	CTACATCTGC	TIGCIGCIGG	GATACCCGTC	ACCACCCCTG	GTCCTTTTAC	300
13	TGTGCCATGG	CAGGCAGTCT	CGGCTTGGGC	CCTCATGGCC	ACACAGACCT	TCTACTCCAC	360
	AGGCCACCAG	CCTGTCTTTC	CAGCCATCCA	TTGGCATGCA	GCCTTCGTGG	GATTCCCAGA	420
20	GGGTCATGGC	TCCTGTACTT	GCTGCCTGC	TTTGCTAGTG	GGAGCCAACA	CCTTTGCCTC	480
	CCACCTCCTC	TTTGCAGTAG	GTTGCCCACT	GCTCCTGCTC	TGGCCTTTCC	TGTGTGAGAG	540
25	TCAAGGGCTG	CGGAAGAGAC	AGCAGCCCCC	AGGGAATGAA	GCTGATGCCA	GAGTCAGACC	600
23	CGAGGAGGAA	GAGGAGCCAC	TGATGGAGAT	GCGGCTCCGG	GATGCGCCTC	AGCACTTCTA	660
	TGCAGCACTG	CTGCAGCTGG	GCCTCAAGTA	CCTCTTTATC	CTTGGTATTC	AGATTCTGGC	720
30	CTGTGCCTTG	GCAGCCTCCA	TCCTTCGCAG	GCATCTCATG	GTCTGGAAAG	TGTTTGCCCC	780
	TAAGTTCATA	TTTGAGGCTG	TGGGCTTCAT	TGTGAGCAGC	GTGGGACTTC	TCCTGGGCAT	840
35	AGCTTTGGTG	ATGAGAGTGG	ATGGTGCTGT	GAGCTCCTGG	TTCAGGCAGC	TATTTCTGGC	900
	CCAGCAGAGG	TAGCCTAGTC	TGTGATTACT	GGCACTTGGC	TACAGAGAGT	GCTGGAGAAC	960
	AGTGTAGCCT	GGCCTGTACA	GGTACTGGAT	GATCTGCAAG	ACAGGCTCAG	CCATACTCTT	1020
40	ACTATCATGC	AGCCAGGGGC	CGCTGACATC	TANGACTTCA	TTATTCWATR	ATTCAGGACC	1080
	ACAGTGGAGT	ATGATCCCTA	ACTCCTGATT	TGGATGCATC	TGAGGGACAA	GGGGKCGGT	1140
45	STCCGAAGTG	GAATAAAATA	GCCGGCCTG	GTGACTTGCA	CCT		1183

(2) INFORMATION FOR SEQ ID NO: 148:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 734 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 148:

GAATTCGGCA GAGTGAAGCA TTAGAATGAT TCCAACACTG CTCTTCTGCA CCATGAGACC 60

50

•	AACCCAGGGC AAGATCCCAT CCCATCACAT CAGCCTACCT CCCTCCTGGC TGCTGGCCAK	120
	GATGTCGCCA GCATTACCTT CCACTGCCTT TCTCCCTGGG AAGCAGCACA GCTGAGACTG	180
5	GGCACCAGGC CACCTCTGTT GGGACCCACA GGAAAGAGTG TGGCAGCAAC TGCMTGGCTG	240
	ACCTTTCTAT CTTCTCTAGG CTCAGGTACT GCTCCTCCAT GCCCATGGYT GGGCCGTGGG	300
	GAGAAGAAGC TCTCATACGC CTTCCCACTC CCTCTGGTTT ATAGGACTTC ACTCCCTAGC	360
10	CAACAGGAGA GGAGGCCTCC TGGGGTTTCC CCRRGGCAGT AGGTCAAACG ACCTCATCAC	420
٠	AGTCTTCCTT CCTCTTCAAG CGTTTCATGT TGAACACAGC TCTCTCCRCT CCCTTGTGAT	480
15	TTCTGAGGGT CACCACTGCC ARCCTCAGGC AACATAGAGA GCCTCCTGTT CTTTCTATGC	540
•	TTGGTCTGAC TGAGCCTAAA GTTGAGAAAA TGGGTGCCAA GGCCAGTGCC AGTGTCTTGG	600
00	GCCCCTTTG GCTCTCCCTC ACTCTCTGAG GCTCCAGCTG GTCCTGGGAC ATGCAGCCAG	660
20	GACTGTGAGT CTGGGCASGT CCAAGGCCTG CACCTTCAAG AAGTGGAATA AATGTGGCCT	720
	TIGCTICTAT TTAA	734
25		
	(2) INFORMATION FOR SEQ ID NO: 149:	
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1405 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 149:	
	GGCACAGTGG ACCCCAGACT CCCTCTCCGC CTTTCTCTGC CTGGGGAGAC CCACTGTGTG	60
40	CATGGCATCA CTGACTCCCA TACCTCTGGC TATCAAAGGT TTCTGCCATG GCCACCCTGG	120
	AAGSAAACCA GAGGGAGGTA GACAGGGAGA TCAGGTCCCT TCTACTCTGG TTCCTGCTCT	180
	GTGAAATTGT CTCAGGCTGG CTGTGTCCAG ARGGTCCCTG GTTCTCTCAR GGATGCCAAA	240
45	TCTACAAGAA TCTCTCCTCT TCCAGTTCCT ATAACCTCTC CTTCCTTTTG TCTCTTTAGA	300
	CCTTGGAGTA GTAGCAGCCA GGTTCTTTCT ATCTCTGGGT TAGTGCATTA TCTCTGGTGG	360
50	CTCCCTTACC CAGGACTTTG GGAATGGTCT TTTTGTAATA CATTCTCCTC AAATAATTCA	420
	ATTITGAGIG TICTGIATGI ATCCIGCIGG GAGGITGITA TATACAAAIC ACIGIGCCCG	480
	TTTAGCAGAG AAGGAGACTG AAGCTCAGGG AGGTTAAGTG TCTTTCTCTA GGTCGTATTG	540
55	TGGAGAAAGT GGCTGACTGG GGACTTGAAT GAGGTCCCTA GTTTCATGCT CGGAGGGCAA	60

AGANGAATGT CCAATTGGCC TGAGATAAGC CTCTGGTAAA ATGTACTGTA CATAATAGGT

AATCAATAAA TGTTGGCTGA TGACAAACAT GTTTTCTTTG TTCATTAGTT ATAGTGATTA

	TGTTCTAAAT AACTCCMACA AGGAARTCAG CACATTTGGA ATATCAWTAT CTTTCCATGA	780
5	TAATATCTTT CCMYGGAAAG AWAATGATAT TCCMAACTGG GAGTGTCCCW AGCARATCTG	840
J	ANTOTOTOTA TTGGCCCTGG GGTGGGCCAG CCCCTTAGAC TCTATGGTCT CATTCTCTTT	900
	GTTTACAAAA TTGAGATAAG GCCTTATTCT CTCCCCACCC CACCCATCCA TATTGTTTTG	960
10	AGAATAAAAT GAGAGGATGT GTGTCAAGGG TGTATTTTGG CAATAGTCTC TGAGCCATTT	1020
	TCTGAGCACC TCCATACTGT TGACACTCAA GTAATATTTC ATCAGCATTC CATTCAGGNT	1080
15	CCTCCCTTAA TGAGGTGTGC GATGTACAAG AGTYGTGAGG TGGCAAAGGA TGGGCTCCTG	1140
13	AGGAAACACT TAGGAAACTG GGCTTTCTGC CATTAAAAGA GACAAACCTT TGTGGTGACC	1200
	TAATTAAAGT TTTTAAAATT CAATTTGGAA AGTTAGCAAG CTAGCTCCTK TCCAGGWAAA	1260
20	ATAAGGAGTC AGTGCATGAC CTAACCGGTC CCGGGCTGCT TGCCATTCCA AACAACTGCA	1320
	GTAAGTTTAT CACNITCTIT CAGGGACTGA GGTTTCCAGG CACAGACTTG GATAAGGAAG	1380
25	GATGTCCTAT GGGGTCACAT TGATG	1405
23		
30 35	(2) INFORMATION FOR SEQ ID NO: 150:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2890 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 150:	
40	TTATATGCTA CAGCTACAGT AATTTCTTCT CCAAGCACAG AGGANCTTTC CCAGGATCAG	60
70	GGGGATCGCG CGTCACTTGA TGCTGCTGAC AGTGGTCGTG GGAGCTGGAC GTCATGCTCA	120
	AGTGGCTCCC ATGATAATAT ACAGACGATC CAGCACCAGA GAAGCTGGGA GACTCTTCCA	180
45	TTCGGGCATA CTCACTTTGA TTATTCAGGG GATCCTGCAG GTTTATGGGC ATCAAGCAGC	240
	CATATGGACC AAATTATGTT TTCTGATCAT AGCACAAAGT ATAACAGGCA AAATCAAAGT	300
50	AGAGAGAGCC TTGAACAAGC CCAGTCCCGA GCAAGCTGGG CGTCTTCCAC AGGTTACTGG	360
JU .	GGAGAAGACT CAGAAGGTGA CACAGGCACA ATAAAGCGGA GGGTGGAAA GGATGTTTCC	420
	ATTGAAGCCG AAAGCAGTAG CCTAACGTCT GTGACTACGG AAGAAACCAA GCCTGTCCCC	480
55	ATGCCTGCCC ACATAGCTGT GGCATCAAGT ACTACAAAGG GGCTCATTGC ACGAAAGGAG	540
	GGCAGGTATC GAGAGCCCCC GCCCACCCCT CCCGGCTACA TTGGAATTCC CATTACTGAC	600
	TTTTCAGAAC CCCACTTCCA TTCAGCCAGG AAACCGCCGG ACTACAACGT GGCCCTTCAG	660

	AGATCGCGGA	TGGTCGCACG	ATCCTCCGAC	ACAGCTGGGC	CTTCATCCGT	ACAGCAGCCA	720
	CATGGGCATC	CCACCAGCAG	CAGGCCTGTG	AACAAACCTC	AGTGGCATAA	AYCGAACGAG	780
5	TCTGACCCGC	CCTCCCCC	YTATCAGTCC	CAAGGGTTTT	CCACCGAGGA	GGATGAAGAT	840
	GAACAAGTTT	CTCCTCTTTG	AGGCACAGAC	TTTTCTGGAA	GCAGAGCGAG	CCACCTGAAA	900
10	GGAGAGCACA	AGAAGACGTC	CTGAGCATTG	GAGCCTTGGA	ACTCACATTC	TGAGGACGGT	960
10	GGACCAGTTT	GCCTCCTTCC	CTGCCTTAAA	AGCAGCATGG	GGSTTCTTCT	CCCCTTCTTC	1020
	CTTTCCCCTT	TGCATGTGAA	ATACTGTGAA	GAAATTGCCC	TGGCACTTTT	CAGACTTTGT	1080
15	TGCTTGAAAT	GCACAGTGCA	GCAATCTTCG	AGCTCCCACT	GTTGCTGCCT	GCCACATCAC	1140
	ACAGTATCAT	TCCAAATTCC	AAGATCATCA	CAACAAGATG	ATTCACTCTG	GCTGCACTTC	1200
20	TCAATGCCTG	GAAGGATTTT	TTTTAATCTT	CCTTTTAGAT	TTCAATCCAG	TCCTAGCACT	1260
20	TGATCTCATT	GGGATAATGA	GAAAAGCTAG	CCATTGAACT	ACTTGGGGCC	TTTAACCCAC	1320
	CAAGGAAGAC	AAAGAAAAC	AATGAAATCC	TTTGAGTACA	GTGCTTGTCC	ACTTGTTTAC	1380
25	AATGTCCTCC	TTTTAAAAAA	AAAAAAATGA	GTTTAAAGAT	TTTGTTCAGA	GAGTAAATAT	1440
	ATATCCATTT	AATGATTACA	GTATTATTTT	AAACCTTAAG	TAGGGTTGCC	AGCCTGGTTT	1500
30	CTGAAAAACC	AAATATGCCG	GACAGGGTGT	GGCCACACCA	AGAAGACGGG	AAGACCTGGC	1560
50	TTGTGACCCT	GGCTTCCCAT	GTCCTTCTGG	TCTCACCCGC	GAAGTGCCCT	ATCCTGGAAG	1620
	TATGAAATGT	TAGCCAATTA	ATACCAAGAC	ACCTCATCTG	CTCCTTCCCC	AGTGGATGGG	1680
35	GTTCTTCTGT	AAAACTGTTT	GCACATGGCC	AGGGGAGGGA	ACTAGGACCC	TTGTGTCCTG	1740
	TCTGAGCCTT	ATGGAGGCAG	GACGCTCTCA	TTGGCGGATG	TGTCCTGCTC	CATTGAGATG	1800
40	GATGGCAAAC	CCCATTITTA	AGTTATATTT	CTTTGATTTT	TGTTAATTTA	GAGGTGTAGG	1860
. 40	TTTTGTTTT	TGTTTTTTT	TTTTTTTTA	AGAGAAACAT	TTATAACTGG	ATAGCATTGC	192
	AGTGAAAGCA	GCTTGGGATG	TTGGAGCTAA	TGCCAGCTGT	TTATACTGCT	CTTTCAAGAC	198
45	AGCCTCCCTT	TATTGAATTG	GCATTAGGGA	ATAAACAAGO	CTTTAAACGI	, GATAAAAGAT	204
	CAAAAACCTG	GTTAGACATG	CCAGCCTTTG	CAAGGCAGGI	TAGTCACCAA	AGACTAACCT	. 210
50	CCAAGTGGCT	TTATGGACGC	TGCATATAGA	GAAGGCCTAA	GTGTAGCAAC	CATCTGCTCA	216
50	CAGCTGCTAT	TAACCCTATA	ATGACTGAAA	TGACCCCTCC	: ACTCTATTT	TGTGTTGTTT	222
	TGCACAGACT	CCGGAAAAGT	GAAGGCTGCC	: AATCTGAGTA	GTACTCAAAT	GTGAGGAACT	228
55	GCTGGTCTTG	GATTTTTTT	CCATTAAATT	CAGCTGATC	TATTGATCAC	TAGATAAACG	234
	TAAATAGCTI	CAAATTTTAA	AAGTGGAATT	GCAGIGTTT	TTCACTGTAT	CAAACAATGT	240
60	CAGTGCTTTA	TTTAATAATT	CTCTTCTGT	TCATGGCATT	TGTCTACTIC	CTTATTACAT	246
60							

	TGTCAATTAT	GCATTTGTAA	TTTTACATGT	AATATGCATT	ATTTGCCAGT	TTTATTATAT	2520
	AGGCTATGGA	CCTCATGTGC	ATATAGAAAG	ACAGAAATCT	AGCTCTACCA	CAAGTTGCAC.	2580
5	AAATGTTATC	TAAGCATTAA	GTAATTGTAG	AACATAGGAC	TGCTAATCTC	AGTTCGCTCT	2640
	GTGATGTCAA	GTGCAGAATG	TACAATTAAC	TGGTGATTTC	CTCATACTTT	TGATACTACT	2700
10	TGTACCTGTA	TGTCTTTTAG	AAAGACATTG	GTGGAGTCTG	TATCCCTTTT	GTATTTTTAA	2760
10	TACAATAATT	GTACATATTG	GTTATATTTT	TGTTGAAGAT	GGTAGAAATG	TACTATGTTT	2820
	ATGCTTCTAC	ATCCAGTTTG	TACAAGCTGG	AAAATAAATA	AATATAACAT	AAAAAAAA	2880
15	ААААААААА						2890

## 20 (2) INFORMATION FOR SEQ ID NO: 151:

25

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2399 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 151:

30	GAACTITITCC ATCTGGCAAA CCGGAAACTC CATCCCCATT AAACCAACTC CCCCTTTTGG	60
*	TTTCCCCCCC AGNGGAATAG AATTTGGACN CCCATATAAA TCCAGGAAAC CACCTAAATT	120
25	CTTTAGTNGT TTGTGTTTGC AAGATCTAAG GTCATGGTAA ACATTAAGTT CTTAAAATTT	180
35	TTGGGAGGGA CCAGTGCACC TCTCCCTCTG AATTGTTCNC CAATTTAAAA TTGGAGTAAG	240
	GTTTTAAAAT GTCTNATTCC ATTGGAAGGG TNTGTTATTT CATTTTGAGC CCAGAGGGGA	300
40	GAGGCACATT TTAAATATCA GAATTAGATT AGCTTTGAGT TTGTACAATT GGGAACATAA	360
	TAGATITICA TAAATTATGT GTGCCTTGTT GGAAGTGTCA ACTGTCTTTA TGTCTGCTTG	420
45	TAAAAGTTTC AAAATATGTT TTCCCTCAAA AAGGCAACGT TACTTCATTT GCTTGAATAT	480
45	TATGATAGGA ATGCTTACTG ATATTACTTG ATAGTCATAT ATAGCCTAGG AAATTTAACA	540
	TATATATAAC TATAGCAGTA TTAATAATGA TAGTTGTACT TCTTTAAAAC ATTAAATTTG	600
50	AGGAAACTTT AATGCTGTCT CGTGTACATT GCTTTACTAC AGTGAGGGGG AATATCCTTT	660
•	AGATTGAGCC TCAATTTACT GGTTAGTAGT ATGTGAACTC TGGTATAAAA ACGTAAACTA	720
	GACAGTAGAG CCGATGAATT AAAATTGTAA ATTGCTACAT TGGCATTTTC TACCTCCTTT	780
. <b>55</b>	TCTGTCAGAG TATTACTTTT TCCAGCATTT ATTCTTATTT GTGAGTAAAG AGGAAATGGG	840
	AACCTGAGGT TAAAATTGAC ATTTTTGTTT CATTGAGAAT TTAAGCAGTA GGTACAGGAG	900
60	AAGTGACTTG TCACATTAAT TTGGTGCCTA AATCTGTAAC TACAAGTTGT GATCGACATG	960

	TACAAAÁTGT	CTAAGAAAGG	TCATATGCTG	AATATTTTAC	TTTTCCTGTA	TAGTCTGCAT	1020
5	GATTTGTTTC	ATAAACCCAG	CTTATTTCCT	CCAAAAAGCA	AAATGGTCCT	GTAATTITTA	1080
3	AAGTAAAATA	AACGTGCCAT	TTTGTCTGCA	ATCTATAATT	TCAGGAAGTT	ATTGRAAGTT	1140
	CTGACTCAGG	GCTTTTTAAC	AGTTCAAGCA	ATTGTCAGTT	ATATTTTGGA	AACTCCATCT	1200
10	GTGTAATTCT	CCAGTGCCTT	GAAAGAATTA	TTAACTTGGC	AACACTATTA	AAACTTTATA	1260
	AAAGATGGTC	TTTAGTGCAC	GTGTATCATT	ATATACACGT	TTTAAAGTCA	TATTGCTTAG	1320
15	CTTGTTAATA	ATGATTCTGC	ATGTGTGCTG	GGTTTGGGTA	ATTCTTTAAA	GGAAGTTTTC	1380
13	TAGATTTGCA	CTTGATGTTT	GTTTTTTAAA	AACTGATTAT	TTATGGCCGT	GACACTGTTA	1440
	CCAGAAAAGT	AATTCTAATT	AAGTTATTAT	GCAAAGTCAT	CTATAAGTAG	CATCTGGGAA	1500
20	GAGGAGATSG	AGGCCACAGT	TIGCTATITI	AGTATGAAAG	GAGGATCTGT	TTGGGAAACA	1560
	TAGATTGTCT	TCCCCTCAAA	TGAGGGGAAA	AAAAAAGACC	CTTTGTTCAA	ATGGATTCTG	1620
25	TTGTAAAAA	TTATTTTTAA	AGGAAATCAC	AAATTGTATG	TCATTCTTAA	TGCTAGTCTT	1680
23	ATAGAATAAA	TCCATAAAAT	TGTTTTTATG	TTCAGTATGT	TTATGTCATT	CTAAATGCAG	1740
	CAAATTCAAT	GATAGCAGTT	CAATTGACTC	ATAGCAGTGT	TTTGTATTTT	TTCTAATTCT	1800
30	TTAGCTTTCA	ATATTGGATT	AAAGTCTTGT	TTGTGAATAT	AGTTTCCGTA	TGGCAAATGA	1860
	TTTCTTGCTT	ATTAGCTTTT	GTTAAAGAAT	GCTTAGTAAG	AGCTAAGCTT	TTAAAAGTAA	1920
35	TGCAAACATT	TATCGTTAAT	AAAACCTATG	GTGTAATATC	ATATAATGCT	TTTCTTTGAT	1980
55	CTTTGGAGAA	TTATTCTTTT	ÄTAGTAGTAT	ACATGAATTT	TĢATTTTAA	AGCATTTAAA	2040
	AACAAATCTC	AATACATTAA	AAAACCTGTT	ATTGTTAAAA	RGGAAATTAC	CATGCCTTTA	2100
40	AGAAACAAGG	ATGTACATCT	TCAATTCAGC	ATRAGTGTCC	ACATCTAGAA	GGCTCTCATT	2160
	GCAGTTGTTT	ACAGTTAAGG	TACCTCTATC	TAAAGGGCCA	AAGAAGCATT	TCATAYTTTA	2220
45	ACACCTCACA	TTCTTTCAGG	ATTAAGACAT	ATGAAAATAG	TCTGAATAGG	ATAAATTTGG	2280
	ATAGGAAGTA	ACTTAACCAG	TCTGGGAAGA	TTCAGGCTTT	ттстаткала	AAGCTTATTC .	2340
	CTCTTCACAA	CTCNGGTGGT	AGGNTTTCAT	TTTTCAAGAG	GGTAGATATT	TTAAAGCCA	2399

## (2) INFORMATION FOR SEQ ID NO: 152:

55 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 802 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

	(XI) SEQUENCE DESCRIPTION. DEG 12 No. 111	
	CGTGCCTGTA GTAAGCTCAT CCCTGCCTTT GAGATGGTGA TGCGTGCCAA GGACAATGTT	60
5	TACCACCTGG ACTGCTTTGC ATGTCAGCTT TGTAATCAGA GATTNTGTGT TGGAGACAAA	120
	TTTTTCCTAA AGAATAACWT GAYCCTTTGC CARACGGACT ACGAGGAAGG TTTAATGAAA	180
	GAAGGTTATG CACCCCMGGT TCGCTGATCT ATCAACATCA CCCCATTAAG AATACAAAGC	240
10 -	ACTACATTCT TTTATCTTTT TTGCTCCACA TGTACATAAG AATTGACACA GGAACCTACT	300
	GAATAGCGTA GATATAGGAA GGCAGGATGG TTATATGGAA TAAAAGGCGG ACTGCATCTG	360
15	TATGTAGTGA AATTGCCCCA GTTCAGAGTT GAATGTTTAT TATTAAAGAA AAAAGTAATG	420
	TACATATGGC TGGATTTTTT TGCTTGCTAT TCGTTTTTGT GTCACTTGGC ATGAGATGTT	480
	TATTTTGGAC TATTGTATAT AATGTATTGT AATATTTGAA GCACAAATGT AATACAGTTT	540
20	TATTGTGTTA CCATTTGTGT TCCATTTGCT YCTTTGTATT GTTGCATTTA GTACAATCAG	600
	TGTTTAAACT TACTGTATAT TTATGCTTTC TGTATTTACC AGCTATTTTA AATGAGCTGT	660
25	AACTTTCTAG TAAAGAATTG AAAAGCAAAT CCTCACTAAA GGATACACAG GATAGGATAA	720
	AGCCAAGTCN CATCAACATT AAAAAATACT AAAANANAAA ACACAAAAAA AAAAAANCCC	780
20	GGGGGGGCC CGGAACCCAT TC	802
30		
35	(2) INFORMATION FOR SEQ ID NO: 153:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH:-461 base pairs	
	(B) TYPE: nucleic acid	
40	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 153:	
		60
45	CTAGGAGCAC CGAGCAGCTT GGCTAAAAGT AAGGGTGTCG TGCTGATGGC CCTGTGCGCA	
	CTGACCCGCG CTCTGCNCTC TCTGAACCTG GCGCCCCCGA CCGTCGCCGC CCCTGCCCCG	
	AGTOTOTTOC COGCOGCOCA GATGATGAAC AATGGCCTCC TCCAACAGCC CTCTGCCTTG	180
50	ATGITGCTCC CCTGCCGCCC AGTTCTTACT TCTGTGGCCC TTAATGCCAA CTTTGTGTCC	240
	TGGAAGAGTC GTACCAAGTA CACCATTACA CCAGTGAAGA TGAGGAAGTC TGGGGGCCGA	300
55	TGGAAGAGTC GTACCAAGTA CACCATTACA CCAGTGAAGA TGAGGAAGTC TGGGGGCCGA GACCACACAG GTGGGAACAA GGACAGGGG ATTTAAGCAG TCAAAAGGAA AAACATGTTA	300 360
55		

## (2) INFORMATION FOR SEQ ID NO: 154:

5 (i) SEQUENCE

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2388 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 154:

	(XI)	DEQUERCE I	DOME TECH	. 026			
	GCCCACGCGT	CCGAAAGCGG	AGAACGCTGG	TGGGCCTGTT	GTGGAGTACG	CTTTGGACTG	. 60
15	AGAAGCATCG	AGGCTATAGG	ACGCAGCTGT	TGCCATGACG	GCCCAGGGGG	GCTGGTGGCT	120
	AACCGAGGCC	GGCGCTTCAA	GTGGGCCATT	GAGCTAAGCG	GGCCTGGAGG	AGGCAGCAGG	180
20	GGTCGAAGTG	ACCGGGGCAG	TGGCCAGGGA	GACTCGCTCT	ACCCAGTCGG	TTACTTGGAC	240
20	AAGCAAGTGC	CTGATACCAG	CGTGCAAGAG	ACAGACCGGA	TCCTGGTGGA	GAAGCGCTGC	300
	TGGGACATCG	CCTTGGGTCC	CCTCAAACAG	ATTCCCATGA	ATCTCTTCAT	CATGTACATG	360
25	GCAGGCAATA	CTATCTCCAT	CTTCCCTACT	ATGATGGTGT	GTATGATGGC	CTGGCGACCC	420
	ATTCAGGCAC	TTATGGCCAT	TTCAGCCACT	TTCAAGATGT	TAGAAAGTTC	AAGCCAGAAG	480
30	TTTCTTCAGG	GTTTGGTCTA	TCTCATTGGG	AACCTGATGG	GTTTGGCATT	GGCTGTTTAC	540
50	AAGTGCCAGT	CCATGGGACT	GTTACCTACA	CATGCATCGG	ATTGGTTAGC	CTTCATTGAG	600
	CCCCTGAGA	GAATGGAGTT	CAGTGGTGGA	GGACTGCTTT	TGTGAACATG	AGAAAGCAGC	660
<b>35</b> .	GCCTGGTCCC	TATGTATTTG	GGTCTTATTT	ACATCCTTCT	TTAAGCCCAG	TGGCTCCTCA	720
	GCATACTCTT	AAACTAATCA	CTTATGTTAA	AAAGAACCAA	AAGACTCTTT	TCTCCATGGT	780
40	GGGTGACAG	GTCCTAGAAG	GACAATGTGC	ATATTACGAC	AAACACAAAG	AAACTATACC	840
<del>-1</del> 0	ATAACCCAAG	GCTGAAAATA	ATGTAGAAAA	CTTTATTTTT	GTTTCCAGTA	CAGAGCAAAA	900
	CAACAACAAA	AAAACATAAC	TATGTAAACA	AGAGAATAAC	TGCTGCTAAA	TCAAGAACTG	960
45	TTGCAGCATC	TCCTTTCAAT	AAATTAAATG	GTTGAGAACA	ATGCATAAAA	AAAGTTGCAC	1020
	AAGTTCCTTA	TTTTCCTTAA	TATTTCACTT	CTATTTAATA	CAAGCTGGGA	CATAAAAATT	1080
50	CTGTTGGGGA	TACCTGGGG	AAGATGTGAG	AAACTAATGC	TGAATTCAGC	TTATACATGA	1140
30	TGAAAAGAAA	AACCAGACAA	AAGGAGCACA	TAAATATGCA	TACAGTGTAA	CTGTTATTAT	1200
	TTTAATACCC	ACGATAAGGG	ATTTTTGTTA	GCATGTTTAG	GGGGAACGAG	GATTGGTGGG	1260
55	ATCCTTGGGG	CCACAGGAAT	CTGAGGCAAC	: GGAAGATATA	TAGAGTGATC	GTCCCCCTGC	1320
	CGAAGGAACC	TGGCAYCTGT	CAAGCAGATO	; CIGCAGTICA	AACTTCAGCT	TTTAAGATAG	1380
60	ATAGCTATTG	AAGGCAGAGG	GTCAGCAGGA	GGATGTGTAT	TTCTAATCTA	CCCTGGTAAA	1440

PCT/US98/11422

	GTCATAGGTA	AGACTCAAAA	GCGGGATCTT	ATTCAAAAGG	CAGGTATTTC	CITIGITITIC	1500
	TGTCTTGAAA	TAGCCCCTTC	CCCTAAGGTG	CATTCTCTCA	AGTTTTCAGT	ATTGCTTTAT	1560
5	TTGCAGTGAT	TAAAAGAGAT	GAGAGACTTT	GGAGACAGAC	AACGTAAGCA	ACACATACAC	1620
	ACATGAAATA	CTCTAGACAG	AGATGAATAT	AAATCTGGCC	TAATAACCAG	TTTTCCATGT	1680
10	AACAGTGATT	TTGTGTTTCG	GGCTGAAGCA	GTGGTTATAT	TAAAAGCCAC	TAATTCCCTT	1740
10	ATCCCTTTAA	AAGATTTTTA	CAATTCTCCA	ACCACAAACA	GCACTTCTAA	AACTAACTTT	1800
	ACTTTCTGCC	CATAATTTGT	TCTACATGGA	ААААААААТ	ATTACTTTGG	CCAGGGGTGT	1860
15	GTGTAAATGT	GGCAGAATTC	CTAGGCAGGC	TGACCTTTAC	AGTATGGGCC	TTTAAGATAC	1920
	TGGATCCTGG	TTGGGCAACA	AGTGTCACGC	CTGAAGTTTC	TGAAAACAAA	TTAGAAGACT	1980
20	GTTGGCTTGG	CTAATCTCGT	AGTTCAGGGC	CAAGTTTCTG	TAGTCAGAAT	GAAGAATAAA	2040
20	ATTGAAAGAA	AAAGGGGGAA	ATGCTTATAC	TTGGCATTAA	GTTGAATGCC	TCAAGTCTTA	2100
	ACTATGGCTT	TGTAGATGAG	GCAAAAGATT	TCTTAGTGGT	AAAATTTCTT	CAACAGGTCA	2160
25	ATGCCAATCT	GTATGCCATT	TTAGTAAAGT	AGGTAAGGAG	AGTAGCCGCT	CAGTAACTTT	2220
	GGCACTAAAG	AAAGAGTGTG	GCTCTAGAAC	TTCCAATCCC	ATTGCTAGAT	GTGCCCTTTA	2280
30	AAAGATGGTC	CAGTGCTTTC	AGGGAAGGAT	GTTTAGCCAG	TTTTCCTAGT	ATTIGTTCCT	2340
30	TAAGATTTTT	TGACCTGTGC	<b>ȚTAATAAGA</b> C	GGACGCGTGG	GTCGACCC		2388

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## (2) INFORMATION FOR SEQ ID NO: 155:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 642 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 155:

45	(AL) BEGOEVED TOTAL TOTA	
45	AAAACAGACC ATTTAAAAAC TCAGACAAGA TTATATTTAA TATATTAATT ACTAAAAAGG	60
	CACAAGATTA CACTGAACAT ATTAGCTACT AAAAAGGCAC TGCTAAGACA TTCAAGCAAA	120
50	TAGCTATTAC ACACTACTGC AGATTTTACA GGTTTCTAAT TCTAACATAT GTTTGAAAAA	180
	TCCGTGAGTA TTCCAAAATA TATTTAATAA TGGAATATCT GCATTAATAT ACCATCCATG	240
E E	TGTTTTTACC ATTTGCCTTA ATATTGAATA TACTGTTTAC CTCACACTAA AAAGAAAACC	300
<b>5</b> 5	AGAAGCCTTA TTTGTGATTT TGGGAGTGGA AGCTTCCATT TTTGTGTCAA AAATGAATCC	360
	TGATTCTTAT GGAAATCTCT GTTATTAAGA TATTTCAAGA TGAGACAACA CTGAAGATCA	420
60	AATTGTGTTT AGTATCACTA TCTTCTCTCC TCGTTTCTCT CTTACTCCTC ATCCTCCCAG	480

WO 98/54963 PCT/US98/11422

	AATCTACCAG TITATGGTAG AAAGATGGGA ACCTTATTTG AATGTGTTTT TTTTTTTCCA	540
5	TGATGTCCAA TTTTGTTGTG GGAAAGGATT TGGATAAAAT TTTTGTTTAA ATTTTGGTAG	600
	ATTTTTATCT ATACAAATTT AAATAAAATT ATGTTTTGTA AG	642
10		•
10	(2) INFORMATION FOR SEQ ID NO: 156:	
15	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 1251 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: double</li> <li>(D) TOPOLOGY: linear</li> </ul>	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 156:	
,	GCCGCTGCCC CTCCACGGAG TTGCTGATCA TCTGGGCTGT GATCCACAAA CCCGGTTCTT	60
	TGTCCCTCCT AATATCAAAC AGTGGATTGC CTTGCTGCAG AGGGGAAACT GCACGTTTAA	120
25	AGAGAAAATA TCACGGGCCG CTTTCCACAA TGCAGTTGCT GTAGTCATCT ACAATAATAA	180
٠.	ATCCAAAGAG GAGCCAGTTA CCATGACTCA TCCAGGCACT GAGCATATTA TTGCTGTCAT	240
30	GATAACAGAA TTGAGGGGTA AGGATATTTT GAGTTATCTG GAGAAAAACA TCTCTGTACA	300
50	AATGACAATA GCTGTTGGAA CTCGAATGCC ACCGAAGAAC TTCAGCCGTG GCTCTCTAGT	360
	CTTCGTGTCA ATATCCTTTA TTGTTTTGAT GATTATTTCT TCAGCATGGC TCATATTCTA	420
35	CTTCATTCAG AAGATCAGGT ACACAAATGC ACGCGACAGG AACCAGCGTC GTCTCGGAGA	480
٠	TGCAGCCAAG AAAGCCATCA GTAAATTGAC AACCAGGACA GTAAAGAAGG GTGACAAGGA	540
40	AACTGACCCA GACTTTGATC ATTGTGCAGT CTGCATAGAG AGCTATAAGC AGAATGATGT	600
40	CGTCCGAATT CTCCCCTGCA AGCATGTTTT CCACAAATCC TGCGTGGATC CCTGGCTTAG	660
	TGAACATTGT ACCTGTCCTA TGTGCAAACT TAATATATTG AAGGCCCTGG GAATTGTGCC	720
45	GAATTTGCCA TGTACTGATA ACGTAGCATT CGATATGGAA AGGCTCACCA GAACCCAAGC	780
	TGTTAACCGA AGATCAGCCC TCGGCGACCT CGCCGGCGAC AACTCCCTTG GCCTTGAGCC	840
	ACTICGAACT TCGGGGATCT CACCTCTTCC TCAGGATGGG GAGCTCACTC CGAGAACAGG	900
50	AGAAATCAAC ATTGCAGTAA CAAAAGAATG GTTTATTATT GCCAGTTTTG GCCTCCTCAG	960
	TGCCCTCACA CTCTGCTACA TGATCATCAG AGCCACAGCT AGCTTGAATG CTAATGAGGT	1020
55	AGAATGGTTT TGAAGAAGAA AAAACCTGCT TTCTGACTGA TTTTGCCTTG AAGGAAAAAA	1080
	GAACCTATTT TIGTGCATCA TTTACCAATC ATGCCACACA AGCATTTATT TTTAGTACAT	1140

TTTATTTTT CATAAAATTG CTAATGCCAA AGCTTTGTAT TAAAAGAAAT AAATAATAAA

ATAAAAAAA AAAAACCCCG GGGGGGGCCC GGTCCCCAAT TGGCCCTATG G 1251

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## (2) INFORMATION FOR SEQ ID NO: 157:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2127 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 157:

.5		
	CCGGCGGGAG AGGGAAGCTG CAGCGAGAGG CGCGGATCTC AGCGCGGGAG CAGTGCTTCT	60
	GCGGCAGGCC CCTGAGGGAG GGAGCTGTCA GCCAGGGAAA ACCGAGAACA CCATCACCAT	120
20	GACAACCAGT CACCAGCCTC AGGACAGATA CAAAGCTGTC TGGCTTATCT TCTTCATGCT	180
	GGGTCTGGGA ACGCTGCTCC CGTGGAATTT TTTCATGACG GCCACTCAGT ATTTCACAAA	240
25	CCGCCTGGAC ATGTCCCAGA ATGTGTCCTT GGTCACTGCT GAACTGAGCA AGGACGCCCA	300
25	GGCGTCAGCG CNCCCTGCAG CACCCTTGCC TGAGCGGAAC TCTCTCAGTG CCATCTTCAA	360
	CAATGTCATG ACCCTATGTG CCATGCTGCC CCTGCTGTTA TTCACCTACC TCAACTCCTT	420
30	CCTGCATCAG AGGATCCCCC AGTCCGTACG GATCCTGGGC AGCCTGGTGG CCATCCTGCT	480
	GGTGTTTCTG ATCACTGCCA TCCTGGTGAA GGTGCAGCTG GATGCTCTGC CCTTCTTTGT	540
25	CATCACCATG ATCAAGATCG TGCTCATTAA TTCATTTGGT GCCATCCTGC AGGGCAGCCT	600
35	GTTTGGTCTG GCTGGCCTTC TGCCTGCCAG CTRACACGGC CCCCATCATG AGTGGCCAGG	660
	GCCTAGCAGG CTTCTTTGCC TCCGTGGCCA TGATCTGCGC TATTGCCAGT GGCTCGGAGC	720
40	TATCAGAAAG TGCCTTCGGC TACTTTATCA CAGCCTGTGC TGTKATCATT TTGACCATCA	780
	TCTGTTACCT GGGCCTGCCC CGCCTGGAAT TCTACCGCTA CTACCAGCAG CTCAAGCTTG	840
45	AAGGACCCGG GGAGCAGGAG ACCAAGTTGG ACCTCATTAG CAAAGGAGAG GAGCCAAGAG	900
45	CAGGCAAAGA GGAATCTGGA GTTTCAGTCT CCAACTCTCA GCCCACCAAT GAAAGCCACT	960
	CTATCAAAGC CATCCTGAAA AATATCTCAG TCCTGGCTTT CTCTGTCTGC TTCATCTTCA	1020
50	CTATCACCAT TGGGATGTTT CCAGCCGTGA CTGTTGAGGT CAAGTCCAGC ATCGCAGGCA	1080
	GCAGCACCTG GGAACGTTAC TTCATTCCTG TGTCCTGTTT CTTGACTTTC AATATCTTTG	1140
	ACTOGTTGGG CCGGAGCCTC ACAGCTGTAT TCATGTGGCC TGGGAAGGAC AGCCGCTGGC	1200
55	TGCCAAGCTG GNTGCTGGCC CGGCTGGTGT TTGTGCCACT GCTGCTGCTG TGCAACATTA	1260
	AGCCCCGCCG CTACCTGACT GTGGTCTTCG AGCACGATGC CTGGTTCATC TTCTTCATGG	1320
60	CTGCCTTTGC CTTCTCCAAC GGCTACCTCG CCAGCCTCTG CATGTGCTTC GGGCCCAAGA	1380

	AAGTGAAGCC	AGCTGAGGCA	GAGACCGCAG	AGCCATCATG	GCCTTCTTCC	TGTGTCTGGG	1440
5	TCTGGCACTG	GGGGCTGTTT	TCTCCTTCCT	GTTCCGGGCA	ATTGTGTGAC	AAAGGATGGA	1500
-	CAGAAGGACT	GCCTGCCTCC	CTCCCTGTCT	GCCTCCTGCC	CCTTCCTTCT	GCCAGGGGTG	1560
	ATCCTGAGTG	GTCTGGCGGT	TTTTTCTTCT	AACTGACTTC	TGCTTTCCAC	GGCGTGTGCT	1620
0	GGCCCGGAT	CTCCAGGCCC	TGGGGAGGGA	GCCTCTGGAC	GGACAGTGGG	GACATTGTGG	1680
	GTTTGGGGCT	CAGAGTCGAG	GGACGGGGTG	TAGCCTCGGC	ATTTGCTTGA	GTTTCTCCAC	1740
15	TCTTGGCTCT	GACTGATCCC	TGCTTGTGCA	GGCCAGTGGA	GGCTCTTGGG	CTTGGAGAAC	1800
IJ	ACGTGTGTCT	CTGTGTATGT	GTCTGTGTGT	CTGCGTCCGT	GTCTGTCAGA	CTGTCTGCCT	1860
	GTCCTGGGGT	GGCTAGGAGC	TGGGTCTGAC	CGTTGTATGG	TTTGACCTGA	TATACTCCAT	1920
20	TCTCCCCTGC	GCCTCCTCCT	CTGTGTTCTC	TCCATGTCCC	CCTCCCAACT	CCCCATGCCC	1980
	AGTTCTTACC	CATCATGCAC	CCTGTACAGT	TGCCACGTTA	CTGCCTTTTT	тааааатата	2040
25	TTTGACAGAA	ACCAGGTGCC	TTCAGAGGCT	CTCTGATTTA	AATAAACCTT	TCTTGTTTTT	. 2100
دے	TTCTCCATGG	АААААААА	AAAAAA				2127

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#### (2) INFORMATION FOR SEQ ID NO: 158:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1625 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 158: 40

CAAAAGATCT ATAATCAGGA CATTGTTTAT GTAAGTTGGA CAANAAAAT TCTTCCCCTT 60 TATGTCCACC CTTCCTATGA TTGCAAGACA AAATTTCCCT CCTTTACCTC ATCCCTATAA 120 CATGGGAGGC TGAGAAAAAT GAGGGGAGAT GGAACCAGAT ACAAGGAGAT CCAATAAGAG 180 AAGCTTATTT AAATATTGTG AAATAAAGGA AGAMCCAAAG CATTTTTTTA AGTGGGGAAT 240 300 CCTTTTGAAC AGTTATTATT TATCCATATT ATTAAYAACA TCTTTTCTGA CAAAATCCAT CAGATGAAGT GTAAATGGAT AATCTTTTAA TGGATCTAAA CCTAGAAAGT TTCACTTACT 360 420 GITCATGTCC GTGTTCCAGA ATTGTGAAAT GGTGTGTGGT TTTGCTTTCC AAGTTCTTCT CTGCCTCCTC TTAATTCTCT AATTCCATGT CTTACAGAAG AATGAGAAAT TTCTTTCTTA 480 CTTGAGTATC ATGCTCTAAA AAACTTGGCT TCAGTCACAG AAACGCTGGC TCTCCTGTGC 540 TTATATTGAA GCCAACTGCC TTTAATTCTT GGGCCCTCTT ATATTTTAA GGTGCAAAAT 600

•	TTGAAGTCTC	AGTCACCAGA	CACAGGTTCT	ATACAATTAA	TGATGAGCTG	GAGAAGTAAT	660
	ATGTAGCTAA	TTTTTCAAAA	GCATTGAATA	TACTTTCCGG	AAAGAAAACA	GAAATTAAAT	720
5	ATTGCCACAT	CTTGCCAGAA	TCCCATCTGA	CACCTTAACT	TTGTCAGGTT	TCCTACAACT	780
	TGCTAATCAA	GTTTTATACA	TTCTAAATCT	CCCCAGTTTC	TTTGGGGCTG	GAAGATGCAA	840
10	CTTCCATTTA	ATAGAAACTT	TGAAATCTTG	GGGTAAGGGA	GCAGTGGGG	GACTAGGGAG	900
10	AAGGATAAGA	AATAGAATTA	TTGAAAAGCC	CCCACCAGGG	ACCTTCCTGG	CCAGAATATG	960
	CAGAGTAATT	CCTGCTGGCT	TCACCTTTGA	AAGTCCCTCG	AAACTATGCA	GATGAAACTG	1020
15	AGTCTGTTTT	TGATATTGTC	AGATGTATTC	TACCTTGGAA	GTCCCNACAC	CTAAACTGGA	1080
	ATTCTTGTAT	TTACATCTCC	TCCACTGTCC	CCCACACCAC	CCCTCAATTC	CTGCTGCCCC	1140
20	TGCTAATGTT	AAGCATTTTT	CTCTTGTTAT	CATCAGGTTC	ACATTAAAAM	CAGRTACTTA	1200
20	CAAACTGACT	TGAAGCACAG	ATACTTTTAC	GAATGTGATA	AAATATTTTC	TTAAGAAAAG	1260
	GAAAGAGGAT	GTGGGTCAAA	TAAAACACCG	CATGGATGTT	GATTGGTGAA	TACTGGTGTA	1320
25	AGAAAAGGGA	GCTCAGGAAT	TTTTATTACT	GTATTTGTAA	ATGAGTTTGA	AGGAATTTGT	1380
	AAATGCCACT	GGTACATTTT	TAAGGTGACA	CATTTGCTCC	TTATAAAGTT	TTAAAAATT	1440
30	ACAGGGTAAG	CTTAAATGAC	GTTTGCCAGT	AGTTTTACTT	TATATAATCA	ATATTGATAT	1500
20	TGTTGCTGAA	CTATGTAACT	TTATGATGCA	TTTTTCAGTC	CCTTTTCAGA	GCAAATGCTT	1560
	TTGCAATGGT	AGTAATGTTT	AGTTTAAATT	GACTTAATAA	ATTMTTACCT	GAGCAAAAAA	1620
35	AAAAA						1625

## 40 (2) INFORMATION FOR SEQ ID NO: 159:

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#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1687 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 159:

50	CGGGGTCACC	AGTTATTAGA	GGAAGTAACA	CAAGGGGATA	TGAGTGCAGC	AGACACATTT	60
	CTGTCCGATC	TGCCAAGGGA	TGATATCTAT	GTGTCAGATG	TTGAGGACGA	CGGTGATGAC	120
55	ACATCTCTGG	ATAGTGACCT	GGATCCAGAG	GAGCTGGCAG	GAGTCAGGGG	ACATCAGGGT	180
	CTAAGGGACC	AAAAGCGTAT	GCGACTTACT	GAACTGCAAG	ATGATAAAGA	ĠGAGGAGGAG	240
	GAGGAGAATC	CACTGCTGGT	ACCACTGGAG	GAAAAGGCAG	TACTGCAGGA	AGAACAAGCC	300
60	AACCTGTGGT	TCTCAAAGGG	CAGCTTTGCT	GGGNATCGAG	GACGATGCCG	ATGAAGGCCC	360

	TGGAGATCAG	TCAGGCCCAG	CTGTTATTTG	AGAACCGGYG	GAAGGGACGG	CAGCAGCAGC	420
5	AGAAGCAGCA	GCTGCCACAG	ACACCCCCTT	CCTGTTTGAA	GACTGAGATA	ATGTCTCCCC	480
J	TGTACCAAGA	TGAAGCCCCT	AAGGNAACAG	AGGCTTCTTC	GGGGACAGAA	GCTGCCACTG	540
	GCCTTGAAGG	GGAAGAAAAG	GATGGCATCT	CAGACAGTGA	TAGĆAGTACT	AGCAKTGAGG	600
10	AAGAAGAGAG	CTGGGAACCC	TCCGTGGTAA	GAAGCGAASC	GTGGGCCTAA	AGTCAGATGA	560
	TGACGGGTTT	GAGATAGTGC	CTATTGAGGA	CCCAGCGAAA	CATCGGATAC	TGGACCCCGA	720
15	AGGCCTTGCT	CTAGGTGCTG	TTATTGCCTC	TTCCAAAAAG	GCCAAGAGAG	ACCTCATAGA	780
13	TAACTCCTTC	AACCGGTACA	CATTTAATGA	GGATGAGGG	GAGCTTCCGG	ACTCCTTTCT	. 340
	GCAAGAGGAA	AAGCAGCACC	GGATACGACA	CTTCCCTCTT	GGTAAGAAGG	AGGTGGAGCA	900
20	TTACCGGAAA	CCCTCCCCCC	AAATCAATGC	ACGTCCCATC	AAGAAGGTGG	CTGAGGCTAA	960
	GGCTAGAAAG	AAAAGGAGGA	TGCTGAAGAG	GCTGGAGCAG	ACCAGGAAGA	AGGCAGAAGC	1020
25	CGTGGTGAAC	ACAGTGGACA	TCTNCAGAAC	GAGAGAAAGT	GGCACAGCTG	CGAAGTCTCT	1080
	ACAAGAAGGC	TGGGCTTGGC	AAGGAGAAAC	GCCATGTCAC	CTACGTTGTA	GCCAAAAAAG	1140
	CTCTCCCCC	CAAAGTGCGC	CGGCCAGCTG	GAGTCAGAGG	TCATTTCAAG	GTGGTGGACT	1200
30	CAAGGATGAA	GAAGGACCAA	AGAGCACAGC	AACGTAAGGA	ACAAAAGAAA	AAACACAAAC	1260
	GGAAGTAAGC	AGAGCTGCCA	GGCTCCCAGG	AGAGCATGGG	GACTAGGAGG	AAGGGTGTGG	1320
35	CATGGCTCAG	TCTGGCCCCC	TTGATTACCG	GCCTAGCCCC	TGCTCACATC	ACAGCTGTCT	1380
	GAAGAACAGT	GAGGTGGAGT	GCCTAGAACT	CCCGTGGTGG	TCCTGAGCAG	AGAGGAGGAT	1440
	GTCCTCCTGC	CTGCCTGAAG	GTCTCCCATG	AAAACACTGC	TGAACTGTGT	TGACACTCAT	1500
<b>40</b> ,	GACCCTTTTT	TTAAACCGTT	AAAGGGAAGT	TCGCTGTTGG	AGCGATACTC	AATGTAGTCA	1560
	GTCTACACCT	GGACGTGTGG	GCCACTTAAG	CCCTCCCCAC	CCCCATCCTA	TTCCTRAATA	1620
45	AAACCAGGAT	AATGGAARAA	ААААААААА	AAAAAAAAAG	GGGGGGCCCN	TAAAGGGNCC	1680
-	CANNTTT						1687

- (2) INFORMATION FOR SEQ ID NO: 160:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 1842 base pairs
      - (B) TYPE: nucleic acid.
      - (C) STRANDEDNESS: double
      - (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 160:

	GGATGACAGA	TIGCGACANA	GATTIGIGAC	CCTTCCTGCT	GAACTTCAGA	GGGAGCTGAA	6
	ANCAGCGTAT	GATCAAAGAC	AAAGGCAGGG	CGAGAACAGC	ACTCACCAGC	AGTCAGCCAG	120
5	CGCATCTGTG	CCCCGAGAAT	CCTTTACTTC	ATCTAAAGGC	AGCAGTGAAA	GAAAAGAAAA	180
	GAAACAAGAA	GAAAAAAACC	ATTGGTTCAC	CAAAAAGGAT	TCAGAGTCCT	TTGAATAACA	240
10	AGCTGCTTAA	CAGTCCTGCA	AAAACTCTGC	CAGGGGCCTG	TGGCAGTCCC	CAGAAGTTAA	. 300
10	TTGATGGGTT	TCTAAAACAT	GAAGGACCTC	CTGCAGAGAA	ACCCCTGGAA	GAACTCTCTG	360
	CTTCTACTTC	AGGTGTGCCA	GGCCTTTCTA	GTTTGCAGTC	TGACCCAGCT	GGCTGTGTGA	420
15	GACCTCCAGC	ACCCAATCTA	GCTGGAGCTG	TTGAATTCAA	TGATGTGAAG	ACCTTGCTCA	480
	GAGAATGGAT	AACTACAATT	TCAGATCCAA	TGGAAGAAGA	CATTCTCCAA	GTTGTGAAAT	540
20	ACTGTACTGA	TCTAATAGAA	GAAAAAGATT	TGGAAAAACT	GGATCTAGTT	АТААААТАСА	600
20	TGAAAAGGCT	GATGCAGCAA	TCGGTGGAAT	CGGTTTGGAA	TATGGCATTT	GACTTTATTC	660
	TTGACAATGT	CCAGGTGGTT	TTACAACAAA	CTTATGGAAG	CACATTAAAA	GTTACATAAA	720
25	TATTACCAGA	GAGCCTGATG	CTCTCTGATA	GCTGTGCCAT	AAGTGCTTGT	GAGGTATTTG	780
	CAAAGTGCAT	GATAGTAATG	CTCGGAGTTT	TTATAATTTT	AAATTTCTTT	TAAAGCAAGT	840
30	GTTTTGTACA	TTTCTTTTCA	AAAAGTGCCA	AATTTGTCAG	TATTGCATGT	AAATAATTGT	900
	GTTAATTATT	TTACTGTAGC	ATAGATTCTA	TTTACAAAAT	GTTTGTTTAT	AAAGTTTTAT	960
	GGATTTTTAC	AGTGAAGTGT	TTACAGTTGT	TTAATAAAGA	ACTGTATGTA	TATTTGGTAC	1020
35	RGGCTCCTTT	TKGTGAAYCC	TTAAAAACTC	AACTCTAGGA	RGCAACTACT	GTTTATTATA	1080
	CTAAARGGCT	GAAAAMCCTC	CAGGCCAGAC	TGCTAAGCTC	TGAAATYCCT	GAGAGGTCTC	1140
40	AGACCGGGAT	TCTACTTGTT	CCAAGAAAGG	GTAAAGCTTC	TAAACCATCT	TATTCTTGTC	1200
	TCCAAGCATG	AACACAGGAG	CATGTYAAGA	AAATCTTTAC	TACTTTCTYC	CATGCGGAGA	1260
	AATCTACATA	TTTTGAATTA	GAAACACCCT	CACACCCACT	TGAAGATTTT	TTTCCTGGGA	1320
45	ACATTATGTC	CCGTAGATCA	GAGGTGGTGT	TETETTTTTG	CTTCTACTGG	CCATTGAGAA	1380
	ACTITGATGA	TAAAAAAGAA	CGGTATAGAT	TTTTCAAACG	TATATAAAAT	ATTTTTATGT	1440
50	TATATGTTAT	GCCATAACTT	TAAAATAAAA	ATAGTTTAAA	ATTCTATGCT	AGTGGATATT	1500
	TGGAACTTTT	TCCTCAAACA	AACACCCCAC	ACTGACTTCA	GCAAAACCCT	AAAACTAGCT	1560
	ACAGATTACT	ACTACGAATG	AATCATYAAG	TTTTGTGTCT	GCAACAATTT	AGAAGCACTA	1620
55	AGCCCAAATA	TCAGGAAATG	TGTGTATGAT	GGAATTTTCT	AGGACAAAAC	AGATCAAGAT	1680
	TAAAACAGGA	TCAAGGATTA	ATGGTATAAA	AATGGTCTAC	TAAAACAGGA	TCAAGGATTA	1740
60	AAACAGGATC	AAGGATTAAT	GGTATAAAAA	TCTCTACTGG	TTACCGGGTG	GCNGGGCCAT	1800

-	ACAGGGTAGT GGTGGATGGA TAGTTTAGTT TGGNAAGGGT AA	1842
5	(2) TITTORIVITYON FOR GEO TO NO. 161.	
	(2) INFORMATION FOR SEQ ID NO: 161:	
10	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 770 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: double</li> <li>(D) TOPOLOGY: linear</li> </ul>	•
15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 161:	
13	GGCACGAGCC CTATGCTGTT CTTGTGATAA TGAGTGAGTC TCACAAGATC TGGTGGTGTT	60
•	ATAGGCATCT GGCATTTCCC CTGCTGACGC TCATTCTCTA TCCTGCCACC CTGGGAAGAA	120
20	GTGTCTTCTG TCATGATTGT AAGTTTCCTG AGGCCTCCCC AGCTATGTAG AACTGTGAGC	180
	CAATTAAACC TCTTTTCTCT ATAAATTATC CAGTCTTATA TATTTCTTCA TAGCAGTGTG	240
25	AGAACAGATA ATACCGTAAA TTGGTATCAC AGAGAGTGGG GTGTTGCTAT AAACACATCT	300
	GAAAATGTTA AAGCAAATTT GGAACTGGGT AACAGGCAAA GGCTGGAACA GTTKGAAGAA	360
	CAGTTAAGAA GAAGACAGGA AAATATGAGA AATCTTGAAA CTTCCTAGAG TCTTAAAGGT	420
30	CTCAGAAGAC ATGAAGATGT GGGAAGCTTT GGAACTTCCT AGAGACTTGT TTGAATGGCT	480
	TTGACCAAAA TGCTGATAGT GATATGGACA ATGAAGTCCA GGCTGAGCTT ATCCAGACAG	540
35	ACATAAGAAG CTCGCTGGGA ACTTGAGTAA AGATCACTCT TGCTAGGCAA AGAGACTGGT	600
55	GGCCTTTTTT CCTCTGCCCT AGAGATCTGT GGAAATCTGA ACCTGAGAGA GATGATTTAG	660
	GGTATCTGGC AGAAGAAATA TCTAAGCGGC AAAACCTTCM AGAGGAAGCA GAGCATAAAC	720
40	GTTTGAAAAA TTTGCAGCCT GACNATGGGA GACCAAAGTT AAACCCAATT	770
45	(2) INFORMATION FOR SEQ ID NO: 162:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 519 base pairs	
50	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 162:	
55	GAATTCGGCA CGAGCTGAGA GGCACAGGAG CAACAGCCAG TGCCCCCTGC AGAGGACCAC	60
55	TGGGGTCACA GACTTCARAC CTGATGACCT GGGCTCAGAT CCCAGCTCTG CACCTACCAG	120
		180
60	CCGTGTGACA AGGTGTCCTC TCTGAGCCTC AGTCACACAC TGCCTTAACG GTTGGGCCTC	190

WO 98/54963 PCT/US98/11422

414

	ATGGAGCTGT	TTGTGAAGGT	TAAATGGGAA	GACATAAAGC	ACTTAGCCCA	GAGCCAAGGA	240
	CATGCTGAAT	AGGATAATGG	TGGCCTCCTT	TGGCGCTGTG	CTGGTGCAGG	TGTGCCGAGG	300
5	AAYTGGGCAG	GGGTGACAGA	TACCTCTTCT	AACCTAGTTC	CTTTCCAAGA	ACCTAATTGG	360
	TGTCTCTCCC	TCCCCCAGGC	AATTGGAAGG	AGGAGGCTGG	GCCCAGCCC	CAGAATACGG	420
10	GAGGTTTCTC	ACCGTGGTAG	GGAAATTGCT	GGCTTGGGGG	TGTGGGCAAC	CACAGTGATC	480
•	GTCTCTCTGC	AGGACGGATG	AGGCTTTGCT	GACAGAGGC			519
	•					•	
15	(2) INFORM	ATION FOR SE	EQ ID NO: 16	53:			
	· (i)	SEQUENCE C	HARACTERIST	ICS:		4	
20		(B) TYP	GTH: 753 ba E: nucleic	acid			
		• - •	ANDEDNESS: OLOGY: line				
25	(xi	) SEQUENCE	DESCRIPTION	: SEQ ID NO	: 163:		
<b>4</b> .5	GGCACGAGCG	GCACGAGCAG	CCAGTTGCTG	ACTGGCACAT	GGCCTCCAGC	GTCCCGGCTG	60
	GTGGGCACAC	TAGAGCCGGA	GGGATCTTCT	TAATTGGTAA	ATTGGATCTT	GAAGCTTCAC	120
30	TGTTTAAATC	TTTTCAGTGG	CTTCCCTTTG	TACTTAGAAA	AAAATGCAAC	TTCTTCTGCT	180
	GGGACTCATC	CGCTCACAGC	CTTCCCCTCC	ACCCTCTCTC	TGCCTCATGC	TCTGCCCCTG	240
35	CCTGCCATGC	CTCCGATACT	CACCTTTTGT	ACCCCAGCAC	CCGTGCCCTC	TGCCCCTCGA	300
	TCTTTGCCTG	GCTGGTTGCT	CCTCACTCAG	TGTTCAGGAC	AAATGCTCCT	GGCCCTACCC	360
40						GCTCTTATTG '	420
40						ATGCAAGCGA	480
	<b></b>					TGTGTGCTCG	540
45		•				TTTGCTAACA	600
						GCAGCCCCA AAGGGTGCAN	660 720
50		CCCGCNAGCG	*		ALICUCATUA	ANGO TO CHIN	720 753
- 0	COLLECTIONIA	CCCCIMICG	WCACCICC10	ogn			, , ,

55 (2) INFORMATION FOR SEQ ID NO: 164:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1400 base pairs

(B) TYPE: nucleic acid

60 (C) STRANDEDNESS: double

## (D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 164:

5	GGCACAGTTT	ATTAATACCT	ATTATGGGAA	AGTCACTITG	GTTGGCATTG	AAAATTACAT	60
	CATCTTTAAA	GCAGTATTTG	TCCCCAGATG	GACTCATCAC	TAGCAAAGAC	TAGGTTCATT	120
10	GGAAGGCATA	GGGTGAGAGA	ATGGGAAGAT	GRAGTGGAGG	CGGGTTGTTA	AAGTGCTGTC	· 180
10	AGTGAGTGAT	TTTGTCTACT	TGAATAATGG	TCCATGTTTG	GGGGCATATT	GTGTTTCATA	240
	AGAAGTGAAA	GGTATTTGCA	AAGTAAGCTA	CAAATGACCC	ATAAATCTGT	TAACAACAGT	300
15	CCTTAATATG	CAAAGATGAA	AAACAAGCAT	TACTGCTACC	CAAAGGGAAC	TGGTGCTTGG	360
	TGATGTGCAG	ATGGGGCTGT	TGGTTAAGAG	AGCTATTACA	GGTTTTCTCT	CTTAGGTTTC	420
20	ATAGGAGGTA	GTTACTGAGA	TGAGATTGTT	TTATCTTTTT	GAATACAGAT	CTCTTGTCTT	480
20	GAGTTAGTTC	TGAGGATGGG	agtaataaag	GAGTTTTTTG	TTTTTTGTT	TGTTTGTTTG	540
	TTTTGGCTCC	TTAGTAATAC	TCCTCTGACA	TTTATTTCTA	TTATTCTTCA	AAGAAAGGAA	600
25	ACCAACTGAA	ATGTTTGCTT	TAACAAACAT	TTTAATAAGT	TCTCTGGGTT	TTTTTTCCC	660
	CTTTTAAAAA	AATTAGCATA	TACCATAGCA	ATAAAAGAAC	TAATGTTAAC	TATTGTATGC	720
30 -	TACAACTTAA	GTGATTTTTC	TAAAGAAGCA	CAATGTCATT	GRAAGTATTA	TTGAAAAGGA	780
50	TCATAGTCAC	ATTGAATTTG	TGAAGGCCAA	AGAAATTGAA	GGGAGTGATA	TTTTCATTTT	840
	ATGATATTCA	CATATTTAGT	AAATTTTGTG	TACAAGAATA	CCAGGCAGAG	TGTTTTACCC	900
35	ATGGAAACAG	GTTTCAGATT	ACTTIGITIT	TACTGTTAGA	GTCTCAAGTT	TAGAAATGCT	960
	AACACTTAAA	TCAGTTTTTT	TCTCACTATA	CTTGAAGATT	GTTAATATTT	TGATATCTTC	1020
40	CTAGCTTGAT	GGAATTTAAA	CATATCTTCA	GATCTGTGAC	AGTGACAGCC	AATAGGACTG	1080
10	ATAATATTAG	CTTCAAACCA	ATAATATCCA	GGGTTAAAAT	AAAAATCATA	GTGAAAGTAC	1140
	GATTGTAAAA	TTATGCTATA	TTAACTITTA	AGTCTGTAAT	AACTTGACAT	CAAAATGTTA	1200
45	TGTAATTACC	ATAAATAATG	GCTAGCGAGA	ACATCTTTGG	AAATTCTCAA	ATTACCTTTC	1260
	TTACTACACT	GTTTGCAGAA	TGAATGTAGA	AATGATCCTG	TTAGCTTTCT	GAATGTTCTG	1320
50	TGGTTGAATG	TGTTTTTGCT	TAAATAAAGC	TTTTGGTATT	TGTTTAAATW	ACAAAAAAAA	1380
50	ааааааааа	AAAAACTCGA					1400

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(2) INFORMATION FOR SEQ ID NO: 165:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2153 base pairs

60 (B) TYPE: nucleic acid

WO 98/54963 PCT/US98/11422

416

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 165: 5 CAGGCCTCAG GGCCTCTGGT GGCTCTGGCC CAGACAGTAT TTGCAGTTCT TGTGCTATGG 60 120 10 CAGGGCTCAA GGGCTGTGGT CCGCTCAGGG TCTCATTTCC CCAGGCCAAG TTCAAGGCAG 180 CAGCCCTTTG TGAGGCGCTC TTGGCCCTGG GCTGGAGGGA GAACTTTAAG CTTTTTTGCT 240 CACAGGACG TGGTATGGGC CCTGGGTGCA GGTGCCCACA TTCTGCTAAT GAGAGCTTTG 300 15 TCTGATCAGT CCTGGGTCCA TCAGTTTGTC CATGTGTCCG GCTGCCAGCC CGTCCCTTGG 360 GATCCTTCCC CTGGGGTGTA GCCTTGTTCA TTAGTATATA CTCATTCCTT CATGCTTTCC 420 20 TCAGCAGAAC ACTTCCACTT CTGAGGTGAG CTTTTGCCCC RTGCCCTTCC TCCACAGGTG 480 TTGCCTTTTT ATAAAGACCT GATAGCAGAA TAAATTGGTG TTTCCCTGTT GACCCAGCAC 540 CATTTCTGTG GGCCTAGAAT ATGGCCCTCA ACCCTTAGAG TGGGGCAGTG AGGGCTTGAG 600 25 GAGTGACCCT TCCTTTCTCA TGGTTTTAGT CATTTTGGCT GCCAGCCCTT AATGGCACAG 660 ATCTGCTGCT TCTAACAGAT GGCCAGGAGG TGACACCGAT TTCAGCCATT GCCAAGGTTA 720 30 GCACCCTCTC CTTTGAGCCT AGGGCCACAC TGTTCATTGT CACTTTAGGC AAGTGCCTGT 780 TTGGCTTTAA AGGTAAGCCT GCCAGCTGTG AGAAGCCTTG GTAACTGATG GACTCATTTC 840 CTGGTCCTTA AAGATGCAGC CTCTTAAGGG CTCCTTGATG GATGCCATCT CTCCTAGCCC 900 35 CCAGCCCTGG TGCCACTGGT GGGCAGGTTC CCATTCTTTG GGGCTGGGAG GGACAGCTTG 960 CCTGTTTCTG GTCACAAATT ACAGTCTTCT CTCCTGTACC ATTCTGTGGC TTCAGCATGG 1020 40 GGGCAGTAGC CTTTCATTAG TGTAGATAGT CATTCCCTGG TAGGGTGGAG GGTAAGACAT AGGGTCTGGA ACTGTTTGGG ACCTTTTGGG GATGTCCTGT GCCTCCCAGA TTCCTMGATT 1140 CTGGGAGGAG AGGCTGCCGC ATTCTGCTGC TCCTCACAGC GAGCAAAGCT GCACCCACTT 1200 45 ACATTCAGTA TTTTCCTGGC ACTACAAAGA GTGGGAAGGC CTGGGATTTG CTGCTGCTCC 1260 CTTAGAGCAG GGCCCCTYTT TTCAGCACTT TGGACACCTG GAGACCCAGC CCTGTTATTT 1320 50 AATGGTAGTG GGCAAGTGTG TGTGCATACT GTCTGCCACT GCTTTCTCCC TGCCCCATGC 1380 CAGAGAGCCC TGTCCCTGCC AGGCCCAGCC TTCTTAGCCC CAACTTGGGA ACAAAGTGCA 1440 ACATGGGATC ATGGGTTGGG GTGCTCAGGT GAGCCCTCTC TATAGTGCTT CCCTGGGCCA 1500 55 AGCTGACACC AGCCCCTGAG GGTGGGGTGG GACGGGTGGT GCTTAAAAGA GGAAGGGGAC 1560 CASTGTAGCA ACTTGCCAGG GACCCCACCC CTCCCTCTCT GGGCCTGTGC AGTGAGCATG 1620

GGGATTCCCA TCAAGGGGCC TGGCACCTGT GCTAGTTACG TAGCCGCTGN TCACGCGCTC

1680

	ACTCCTGACC	ACATGCACGT	TCCCTAGATG	CAGACTGCTT	TGAACTTTAA	AGCTGTACAA	1740
5	TTTGGTTATG	TTTGTGCTGA	CTTAAAATAT	ATTTTAATGA	GGAAAAAATA	ATGGAGAACC	1800
•	CTGGGAAGGA	CCTGGTTCTT	TTGCTTCTCG	GGGAACTGTA	AGCCCTCGCG	TTCTGGGAAT	1860
	CGCTCTCTGC	TGCTCTTTCC	TGGAAGCTAA	GCCTGTCTCC	ACCGCCCGAG	GCCTGCGCCG	1920
10	GTGCTCCCGC	CGCAGTTGCG	TTTGCTTTGG	ACCTTGCGTG	CGGGGGAGGG	GGTGCTCGGT	1980
	CCGAGCCCGC	TCCTTTCTGT	ACACCTAGCG	CTGCCCGCCC	CGCTTGTGTC	TGAGGTCGTG	2040
15	TATGTCAAAA	ATAAAGCCGC	TAGAAACGGA	АААААААА	АААААААА	ААААААААА	2100
10	AAACTCGAGG	GGGGGCCCGT	ACCCAATTAA	CCCNNTATGA	TCTATAAAGC	GTC	2153

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#### (2) INFORMATION FOR SEQ ID NO: 166:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1251 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 166:

GCCCACGCGT CCGCCCACGC GTCCGGCGGT GCGGAGTATG GGGCGCTGAT GGCCATGGAG 60 GGCTACTGGC GCTTCCTGGC GCTGCTGGGG TCGGCACTGC TCGTCGGCTT CCTGTCGGTG 120 35 ATCTTCGCCC TCGTCTGGGT CCTCCACTAC CGAGAGGGGC TTGGCTGGGA TGGGAGCGCA 180 CTAGAGTTTA ACTGGCACCC AGTGCTCATG GTCACCGGCT TCGTCTTCAT CCAGGGCATC 240 GCCATCATCG TCTACAGACT GCCGTGGACC TGGAAATGCA GCAAGCTCCT GATGAAATCC 300 40 ATCCATGCAG GGTTAAATGC AGTTGCTGCC ATTCTTGCAA TTATCTCTGT GGTGGCCGTG 360 TTTGAGAACC ACAATGTTAA CAATATAGCC AATATGTACA GTCTGCACAG CTGGGTTGGA 420 45 CTGATAGCTG TCATATGCTA TTTGTTACAG CTTCTTCAG GTTTTTCAGT CTTTCTGCTT 480 CCATGGGCTC CGCTTTCTCT CCGAGCATTT CTCATGCCCA TACATGTTTA TTCTGGAATT 540 GTCATCTTTG GAACAGTGAT TGCAACAGCA CTTATGGGAT TGACAGAGAA ACTGATTTTT 600 50 TCCCTGAGAG ATCCTGCATA CAGTACATTC CCGCCAGAAG GTGTTTTCGT AAATACGCTT 660 GGCCTTCTGA TCCTGGTGTT CGGGGCCCTC ATTTTTTGGA TAGTCACCAG ACCGCAATGG 720 55 780 AAACGTCCTA AGGAGCCAAA TTCTACCATT CTTCATCCAA ATGGAGGCAC TGAACAGGGA GCAAGAGGTT CCATGCCAGC CTACTCTGGC AACAACATGG ACAAATCAGA TTCAGAGTTA 840 AACAGTGAAG TAGCAGCAAG GAAAAGAAAC TTAGCTCTGG ATGAGGCTGG GCAGAGATCT 900 60

882

	ACCATGTAAA ATGTTGTAGA GATAGAGCCA TATAACGTCA CGTTTCAAAA CTAGCTCTAC	960
	AGTTTTGCTT CTCCTATTAG CCATATGATA ATTGGGCTAT GTAGTATCAA TATTTACTTT	1020
5	AATCACAAAG GATGGTTTCT TGAAATAATT TGTATTGATT GAGGCCTATG AACTGACCTG	1080
	AATTGGAAAG GATGTGATTA ATATAAATAA TAGCAGATAT AAATTGTGGT TATGTTACCT	1140
10	TTATCTTGTT GAGGACCACA ACATTAGCAC GGTGCCTTGT GCAKAATAGA TACTCAATAT	1200
	GTGAATATGT GTCTACTAGT AGTTAATTGG ATAAACTGGC AGCATCCCTG A	1251
15	(2) INFORMATION FOR SEQ ID NO: 167:	
20	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 882 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 167:	
	GACSMTCTAG AACTATGGTC CCCCGGGACT GCAGGAATTC GGCACAGCGG CTGCGGGCGC	60
	GAGGTGAGGG GCGCGAGGTT CCCAGCAGGA TGCCCCGGCT CTGCAGGAAG CTGAAGTGAG	120
30	AGGCCCGGAG AGGGCCCAGC CCGCCCGGGG CAGGATGACC AAGGCCCGGC TGTTCCGGCT	180
	GTGGCTGGTG CTGGGGTCGG TGTTCATGAT CCTGCTGATC ATCGTGTACT GGGACAGCGC	240
35	AGGCGCCGCG CACTTCTACT TGCACACGTC CTTCTCTAGG CCGCACACGG GGCCGCCGCT	300
	GCCCACGCCC GGGCCGGACA GGGACAGGGA GCTCACGGCC GAYTCCGATG TCGACGAKTT	360
	TCTGGACAAK TTTCTCAGTG CTGGCGTGAA GCAGAGTGAC YTTCCCAGAA AGGAGACGGA	420
40	GCAGCCGCCT GCGCCGGGA GCATGGAGGA GAGCGTGAGA RGCTACGACT GGTCCCCGCG	480
	CGAMCCCCGG CGCACCCAGA CCAGGGCCGG CAGCARGCGG ANCGGAGGAR CGTGCTGCGG	540
45	GGCTTCTGCG CCAAYTCCAG CCTGGCCTTC CCCACCAAGG AGCGCGCATT CRACGACATC	600
	CCCAACTCGG AGCTGAGCCA CCTGATCGTG GACGACCGGC ACGGGGCCAT CTACTGCTAC	660
	GTGCCCAAGG TGGCCTGCAC CAACTGGAAG CGCGTRATGA TCGTGCTGAG CGGAAGCTGT	720
50	GCACCGCGTG CGCCTACCGC GACCCGYTGC GNTCCCGCGC GAGCACGTGC ACAACGCCAG	780

CGCGCACTGA CTTCAACAAT TCTGGCGCCG CTACGGGAAG TCTCCCCCAC CTCATGAAGT

CAAGCTCAAG AATACACCAA TTCTTTCTGC GCGACCCTTC TG

<sup>(2)</sup> INFORMATION FOR SEQ ID NO: 168:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1208 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 168:

10	GGGAAACTCA	AAAGGATGAT	GGAATGGTTG	ATGGAGCCAG	AGCCTAGAAG	TRAAGGGATA	. 60
10	CAGAGTGAAG	ATAGAGGTAT	TTACGTATAT	TTWAATATTA	GCTTTGGAAT	TACGTAGGGA	120
	TTCTTAAGAA	AAGATCATGA	CAGGACAGCC	ACATTTGGTA	AAATGTCAGG	GCAGCCAGTG	180
15	CATGGTCCTC	CTGGGGCTCC	TCAGTTGACG	GGTTTAAATC	ATTTCCTGAT	CCCCTGCCC	240
	TGGTTTGAGG	AATGCATACA	GTACGTGAAA	TGCCTGTGGT	ATGAGTTGCA	ATGGGCAATC	300
20	AACCTGGGTA	AATCCAAGAT	TAATGATTAG	TTCTAAAGAT	CCAGTTGAAG	TTCTAGAGTG	360
20	GGAATTTTCC	GTCAAGCARC	TCAGCACAGC	TTTATGCCTG	TTCCTCTAAT	AACGATAGGT	420
	AACAAATAGC	TGTGTKTWCA	CAGCTAGGAR	GATAACCAAA	TCTAGAGTTC	TTGARTCTCA	480
25	TTTAATAAAT	AAKTATTATG	AGTACCAACT	GCATATTTCA	GGCACTGCAT	TTGACTCTGT	540
	TAAATACTGA	TYCCTTAKGA	CMSCCACWIC	AGAWAACMIT	AATCTGTCTG	ATCAATAAAC	600
30	AGCTTGACTT	AGAGRGGTAA	AATAGCTTGC	CACAGGTWAC	CCAATTAGTA	GGTAACAGCG	660
	ACAGAATAAC	AGTGCAGTTA	AAATCTTAGA	CTGGAGACTA	ATTGCATAAG	TTTGAATTTC	720
·	AGTTCTGCTA	TGTAAATTTG	GGTGAGTACC	TTAATTYACC	TGAGTCTCGG	TCTTTATATC	780
35	TGTAGAATGG	AGCTAATGAT	ATTACTTAAT	TTGCTTTATG	TGAGATTAAA	TGTACTAATA	840
	TATGTAAATC	ACTTACAACA	GCAŢŢŢĠĀĊĀ	TATTTGACAT	ACTTAATATA	TTTGCTACTA	900
40	ATACTATTAG	CAACAGCATT	CTGATTTTCC	AAGTTGAAAT	TCAGTGTTTT	CTTTTTTACT	960
	TTGCCATAAT	TTACAATGTT	GTGCTCTGTA	AACCATAAAT	TTCCCTGAGG	TGTTGTCAGG	1020
	TTAAAAAAAA	ATCACTATGG	CCCCARNIMA	CTTGGAAAAT	AGAAATGAGA	CCAGCTTCAT	1080
45	CTATATTCTT	TACTGCAAAT	AACTTAGAAT	TGTAATAGGC	TAATATGTAC	TGGGACTTCC	1140
	AATTTGGGAA	TATGACAAAA	ATAATACTAT	TTAGCTAAAA	CATATACAGA	ACTTATTTTT	1200
50	CCTCTGAA						1208

(2) INFORMATION FOR SEQ ID NO: 169:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1307 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

60 (D) TOPOLOGY: linear

(xi) SEQUE	INCE DESC	RIPTION:	SEO	ID	NO:	169:
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5	GGCACGAGAG	AAAAGAGGTT	GAGAATGTTT	TCTAGCAGGC	AGAATGTGCA	TACATGTTTT	6
	CATGARTGTC	CTTTGGGTGC	TGTTTCTTTT	AAATCCTCTG	TGCACAGGGC	TCTGGCCTTT	120
	ARTAAACTGT	TTTTCTCTCT	TACGTCATGC	TGACTGGGTG	CTAGGGGCTG	ATTACAAAGG	180
10	GGAAGAGTTG	AACAGACATC	AGGGGCCGAT	GAAACCAAAG	GACTAGGAGT	CAGGAGAACA	24(
	AGTCAGGGAT	TAGGAGACAG	CGGTTTGGTT	TATTGTTATC	CAGCTGGAGG	ACTCCTAGGG	300
15	GCAGCAGCAG	GAGGAATACC	AGGGCCACGG	AGGGGCAGGA	GTCTCACAGT	GGAGGGCAGA	360
13	CTCTAACAGA	TGCCAGCTGA	ACGCTCGCTG	GCCCTGGATG	TCATACGAGT	TGGGGACCAG	420
	AAATCTGGGC	TCAGAGAACC	CGTCCAGGGA	GATTTGAAGC	CATGGGTTAT	CTTCTAGAGT	480
20	TGATACTGAT	AATATATTTT	AATTTTTATT	GATGTTTAAT	ACCTTCTGAA	ACAGGAGGGT	540
	AAGATCAGAT	GGGAAGCCCY	TCTGTTGAAG	GATCTTGGGA	ACCTTGGTGG	TTTTTTTTT	600
25	TIGGITITIT	TTTTTTTGAT	CGAGCTGTGG	ACATCCTTCT	TAATTCGATT	NTGAGGATTT	660
23	GTTTAACTAA	AAAGTTCCCA	AACACAGAAA	GGCCTCCCC	ACCTGCTTTG	GGGAGCTGTC	720
	TGTSCTGGGA	GTGCCAGGCA	TCCSATGGGA	CCCATCACTG	CCAGTGTCTG	TGCCTCCCAG	780
30	AGGTCAGCCC	TGTGTCTGCC	CTGGCTCTGT	CTCCTCTGTG	ACAGGGCAGA	GCATTTCTGG	840
	TCAGTTTCTC	CATGGTGCCT	CCCACCCCTT	TGTAAAGTGG	ATGGACATGA	TGGAATTCAG	900
35	TTGTCTCACC	CTGATAGCCT	GGGTGTTGAT	ATTCACTTTA	CCCGCACTCA	GACACAGGCG	960
,,	ACCTTGAAGC	AGTTCTCGGT	GTGTAGAGTC	CACGTGACAG	TCCCCACAGC	CTCCCCAGAT	1020
	AGCTGTGTGC	CTGTGCGCTA	CTCCTGTGCC	ATTTTCCCAA	CTTNGGCGTT	TCACTAAATG	1080
40	CAGCTGATCT	CTCTCTCTGT	GCACTCGTGA	TCCATGTTGA	ACAATACATG	TAGGTTCTTT	1140
	TTCCACGCAA	TGTAAGAACA	TGATATACTG	TACGTTGGAA	AGCATTTACC	TTATTTATAT	1200
15	ACCTGAATGT	TCCTACTACA	CAAATAAACA	TATATTAAAT	WCTAAAAAA	АААААААА	1260
<del>,</del> ,	CTGGAGGGG	GGCCCGGTAC	CCAAATCGCC	GGATAGTGAT	CGTAAAC		1307

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## (2) INFORMATION FOR SEQ ID NO: 170:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1624 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 170:

•	GGCACGAGGT	ceccecece	GCCGCCTGGA	ATTGTGGGAG	TIGIGICIGC	CACTCGGCTG	60
	CCGGAGGCGA	AGGTCCCTGA	CTATGGCTCC	CCAGAGCCTG	CCTTCATCTA	GGATGGCTCC	120
5	TCTGGGCATG	CTGCTTGGGC	TGCTGATGGC	CGCCTGCTTC	ACCTTCTGCC	TCAGTCATCA	180
	GAACCTGAAG	GAGTTTGCCC	TGACCAACCC	AGAGAAGAGC	AGCACCAAAG	AAACRGAGAG	240
10	AAAAGAAACC	AAAGCCGAGG	AGGAGCTGGA	TGCCGAAGTC	CTGGAGGTGT	TCCACCCGAC	300
10	GCATGAGTGG	CAGGCCCTTC	AGCCAGGGCA	GCTGTCCCT	GCAGGATCCC	ACGTACGGCT	360
	GAATCTTCAG	ACTGGGGAAA	GAGAGGCAAA	ACTCCAATAT	GAGGACAAGT	TCCGAAATAA	420
15	TTTGAAAGGC	AAAAGGCTGG	ATATCAACAC	CAACACCTAC	ACATCTCAGG	ATCTCAAGAG	480
	TGCACTGGCA	AAATTCAAGG	AGGGGGCAGA	GATGGAGAGT	TCAAAGGAAG	ACAAGGCAAG	540
20	GCAGGCTGAG	GTAAAGCGGC	TCTTCCGCCC	CATTGAGGAA	CTGAAGAAAG	ACTITGATGA	600
20	GCTGAATGTT	GTCATTGAGA	ĊTGACATGCA	GATCATGGTA	CGGCTGATCA	ACAAGTTCAA	660
	TAGTTCCAGC	TCCAGTTTGG	AAGAGAAGAT	TGCTGCGCTC	TTTGATCTTG	AATATTATGT	720
25	CCATCAGATG	GACAATGCGC	AGGACCTGCT	TTCCTTTGGT	GGTCTTCAAG	TGGTGATCAA	780
	TGGGCTGAAC	AGCACAGAGC	CCCTCGTGAA	GGAGTATGCT	GCGTTTGTGC	TGGGCGCTGC	840
30	CTTTTCCAGC	AACCCCAAGG	TCCAGGTGGA	GGCCATCGAA	GGGGGAGCCC	TGCAGAAGCT	900
	GCTGGTCATC	CTGGCCACGG	AGCAGCCGCT	CACTGCAAAG	AAGAAGGTCC	TGTTTGCACT	960
	GTGCTCCCTG	CTGCGCCACT	TCCCCTATGC	CCAGCGGCAG	TTCCTGAAGC	TCGGGGGGCT	1020
35	GCAGGTCCTG	AGGACCCTGG	TGCAGGAGAA	GGGCACGGAG	CTCCTCCCCG	TGCGCGTGGT	1080
	CACACTGCTC	TACGACCTGG	TCACGGAGAA	GATGTTCGCC	GAGGAGGAGG	CTGAGCTGAC	1140
40	CCAGGAGATG	TCCCCAGAGA	AGCTGCAGCA	GTATCGCCAG	GTACACCTCC	TGCCAGGCCT	1200
,,	GTGGGAACAG	GCTGCTGCG	AGATCACGGC	CCACCTCCTG	GCGCTGCCCG	AGCATGATGC	1260
	CCGTGAGAAG	GTGCTGCAGA	CACTGGGCGT	CCTCCTGACC	ACCTGCCGGG	ACCGCTACCG	1320
45	TCAGGACCCC	CAGCTCGGCA	GGACACTGGC	CAGCCTGCAG	GCTGAGTACC	AGGTGCTGGC	1380
	CAGCCTGGAG	CTGCAGGATG	GTGAGGACGA	GGGCTACTTC	CAGGAGCTGC	TGGGCTCTGT	1440
50	CAACAGCTTG	CTGAAGGAGC	TGAGATGAGG	CCCCACACCA	GGACTGGACT	GGGATGCCGC	1500
<b>J</b> 0	TAGTGAGGCT	GAGGGGTGCC	AGCGTGGGTG	GGCTTCTCAG	GCAGGAGGAC	ATCTTGGCAG	1560
	TGCTGGCTTG	GCCATTAAAT	GGAAACCTGA	AGGCCAAAAA	ааааааааа	АААААААА	1620
55	AAAA						1624

WO 98/54963

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2003 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 171:

10	GGCACGAGCC	AGCTTGCAGG	AGGAATCGGT	GAGGTCCTGT	CCTGAGGCTG	CTCTCCGGG	60
	CCGGTGGCTG	CCCTCAAGGT	CCCTTCCCTA	GCTGCTGCGG	TTGCCATTGC	TTCTTGCCTG	120
15	TTCTGGCATC	AGGCACCTGG	ATTGAGTTGC	ACAGCTTTGC	TTTATCCGCG	CTTGTGTGCA	180
13	GGCCCGGCT	GGGCTCCCCA	TCTGCACATC	CTGAGGACAG	AAAAAGCTGG	GTCTTGCTGT	240
	GCCCTCCCAG	GCTTAGTGTT	CCCTCCCTCA	AAGACTGACA	GCCATCGTTC	TGCACGGGGC	300
20	TTTCTGCATG	TGACGCCAGC	TAAGCATAGT	AAGAAGTCCA	GCCTAGGAAG	GGAAGGATTT	360
	TGGAGGTAGG	TGGCTTTGGT	GACACACTCA	CTTCTTTCTC	AGCCTCCAGG	ACACTATGGC	420
25	CTGTTTTAAG	AGACATCTTA	TTTTTCTAAA	GGTGAATTCT	CAGATGATAG	GTGAACCTGA	480
23	GTTGCAGATA	TACCAACTTC	TGCTTGTATT	TCTTAAATGA	CAAAGATTAC	CTAGCTAAGA	540
	AACTTCCTAG	GGAACTAGGG	AACCTATGIG	TTCCCTCAGT	GTGGTTTCCT	GAAGCCAGTG	600
30	ATATGGGGGT	TAGGATAGGA	AGAACTTTCT	CGGTAATGAT	AAGGAGAATC	TCTTGTTTCC	660
	TCCCACCTGT	GTTGTAAAGA	TAAACTGACG	ATATACAGGC	ACATTATGTA	AACATACACA	720
35	CGCAATGAAA	CCGAAGCTTG	GCGCCTGGG	CGTGGTCTTG	CAAAATGCTT	CCAAAGCCAC	780
33	CTTAGCCTGT	TCTATTCAGC	GGCAACCCCA	AAGCACCTGT	TAAGACTCCT	GACCCCCAAG	840
	TGGCATGCAG	CCCCCATGCC	CACCGGGACC	TGGTCAGCAC	AGATCTTGAT	GACTTCCCTT	900
40	TCTAGGGCAG	ACTGGGAGGG	TATCCAGGAA	TCGGCCCCTG	CCCCACGGGC	GTTTTCATGC	960
	TGTACAGTGA	CCTAAAGTTG	GTAAGATGTC	ATAATGGACC	AGTCCATGTG	ATTTCAGTAT	1020
15	ATACAACTCC	ACCAGACCCC	TCCAACCCAT	ATAACACCCC	ACCCCTGTTC	GCTTCCTGTA	1080
45	TGGTGATATC	ATATGTAACA	TITACTCCTG	TTTCTGCTGA	TIGITITIT	AATGTTTTGG	1140
	TTTGTTTTTG	ACATCAGCTG	TAATCATTCC	TGTGCTGTGT	TTTTTATTAC	CCTTGGTAGG	1200
50	TATTAGACTT	GCACTTTTT	AAAAAAAGGT	TTCTGCATCG	TGGAAGCATT	TGACCCAGAG	1260
	TGGAACGCGT	GGCCTATGCA	GGTGGATTCC	TTCAGGTCTT	TÇCTTTGGTT	CTTTGAGCAT	1320
<b>5 5</b>	CTTTGCTTTC	ATTCGTCTCC	CGTCTTTGGT	TCTCCAGTTC	AAATTATTGC	AAAGTAAAGG	1380
55	ATCTTTGAGT	AGGTTCGGTC	TGAAAGGTGT	GGCCTTTATA	TTTGATCCAC	ACACGTTGGT	1440
	CTTTTAACCG	TGCTGAGCAG	AAAACAAAAC	AGGTTAAGAA	GAGCCGGGTG	GCAGCTGACA	1500
60	GAGGAAGCCG	CTCAAATACC	TTCACAATAA	ATAGTGGCAA	татататата	GTTTAAGAAG	1560

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•	GCTCTCCATT TGGCATCGTT TAATTTATAT GTTATGTTCT AAGCACAGCT CTCTTCTCCT	1620
5	ATTITCATCC TGCAAGCAAC TCAAAATATT TAAAATAAAG TTTACATTGT AGTTATTTTC	1680
,	AAATCTTTGC TTGATAAGTA TTAAGAAATA TTGGACTTGC TGCCGTAATT TAAAGCTCTG	1740
	TTGATTTTGT TTCCGTTTGG ATTTTTGGGG GAGGGGAGCA CTGTGTTTAT GCTGGAATAT	. 1800
10	GAAGTCTGAG ACCTTCCGGT GCTGGGAACA CACAAGAGTT GTTGAAAGTT GACAAGCAGA	1860
	CTGCGCATGT CTCTGATGCT TTGTATCATT CTTGAGCAAT CGCTCGGTCC GTGGACAATA	1920
15	AACAGTATTA TCAAAGAGAA AAAAAAAAAA AAAAAACTCG NGGGGGGCC CGGTACCCAA	1980
10	TTCGCCCTAT AGTGAGCCNA TTC	2003
20	(2) INFORMATION FOR SEQ ID NO: 172:	
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 786 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 172:	
30	GGCACAGCGG CACGAGAAGA CTTTGGTGTT TAAGAGATTA ATGTGTTAGC CAGAACAACT	60
	CATTTCTCTA CCMGTGTGTA GTCCATTTAT CTTTAAAGAT TTTCTATTGG AATAATTTTG	120
35	AAATTACTTT CTTAGTTTTC TTCATTAAAA ACTAAGAAAA TGCTTTGTTT ATTATGAATT	180
	GCTATTTCTC TTGATTATTA TTCTTGGAGA AAGTCTATCA GACGTAATTC TTCTGATTTG	240
40	CTTCTAGGCT AGAGGAAAAT GTGAAAGATG ACAAATGAAA ATTTCAAAGG TTGTCAGTAG	300
	TATGACTTCT TTTATCGTTT GTCATTATCA CAAATATATC AACATAGGAC TTTTAAAAGA	360
	TATTITGTAC ATATTGGGCC TTAGTAGGAT TITGCATGAA TITTITTTTT CITTTATGCC	420

CAGAGAGAAA GAGCAAAGAA ATAACCAAGG GTGATGTACT CGTATTGAAG GTTTACCAAA

TAAGGACTGC TTTTATTATG AACTATAGTC TATATTCTAA GTAAATCAAT TTTTCTATTA

TGTGTTTTTT GTTCCTGCAG GCAAGATCTC TGAACTTTAT GCAGAGGGTT CTTTTAAAAA

AACAAAGTTG AATTTTTTTA TTTCTTGGAA TATTTTTTTT CATTGATTTC TCCCAAGTAG

AGCAGATTCA AATCTCCTTT GTACCCTATG TCTTTTTTGT TTTGCTATTA GCTCAGTATT

CCGTTTCTAC ATTTTCCTTT CCTAGAACCA GTCAATAAAT GACAAAAAAA AAAAAAAAA

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ACTCGA

## (2) INFORMATION FOR SEQ ID NO: 173:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1758 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 173:	
	GGGACGAGCC CTGCCCACCT CCTGCAGCCT CCTGCGCCCC GCCGAGCTGG CGGATGGAGC	60
15	TGCGCACGGG GAGCGTGGGC AGCCAGGCGG TGGCGCGGAG GATGGATGGG GACAGCCGAG	120
15	ATGGCGGCGG CGGCAAGGAC GCCACCGGGT CGGAGGACTA CGAGAACCTG CCGACTAGCG	180
,	CCTCCGTGTC CACCCACATG ACAGCAGGAG CGATGGCCGG GATCCTGGAG CACTCGGTCA	240
20	TGTACCCGGT GGACTCGGTG AAGACACGAA TGCAGAGTTT GAGTCCAGAT CCCAAAGCCC	300
	AGTACACAAG TATCTACGGA GCCCTCAAGA AAATCATGCG GACCGAAGCT TCTGGAGGCC	360
05	CTTGCGAGGC GTCAACGTCA TGATCATGGG TGCAGGGCCR GCCCATGCCA TGTATTTTGC	420
25	CTGCTATGAA AACATGAAAA GGACTTTAAA TGACGTTTTC CACCACCAAG GAAACAGCCA	480
	CCTAGCCAAC GGTATTTTGA AAGCGTTTGT CTGGAGTTAG AAAGTTCTCT TCTTCAACAC	540
30	GTCCCTCCCC AGGGTGTTCC TCCCTGTGAC CCAGCCGCCT CGACTTCGGC CCGCTTGCTC	600
	ACGAATAAAG AACTCAGAGT TGTGTGCA ATGCACACCC AGACACACGC ACGCACACAC	660
25	ACGCGCGCGC ACACACATGC TTTTTTCTGT TCCCCTCCGC TTTCTGAAGC CTGGGGAGAA	720
35	ATCAGTGACA GAGGTGTTTT GGTTTTATTG TTATGTGGGT TTTCTTTTGT ATTTTTTTTG	780
	TTTGTTTTGT TTTTAAACAT TCAAAAGCAA TTAATGATCA GACATAGGAG AAACCCTGAA	840
40	TAGAAACAAA ACTTTTGAAT GCTGGATTCA AAAAAAAAA AAAGTTATCT GGACAGCTTC	900
	TTTGAGACTA TTTAAAAACT GGTACAACAG GTCTCTACAA CGCCAAGATC TAACTAAGCT	960
15	TTAAAAGGTC AAGAAGTTTT ATGGCTGACA AAGGACTCGC GCAACGCAGA AGGCCTTTCC	1020
45	CACCTTAAGC TTCCGGGGAT CTGGGAATTT TACCCCCATT CTCTTCTGTT TGTCTGAGTC	1080
	TCATCTCTCT GCAAGCAAGG GCTGAAATCA TTTTGTTTGG TTGTTTTGAG GGAGAGAGGCC	1140
50	GGGGTGGGGG GGTGCAAATC TGCCAGCAGC TCTTACGTAA GGCATGTTTT ATTGGGGAGG	1200
	GCTGAGCTTT TATTTTCTCC TCTCCAGTGG GGTTGGCTTT TATTGTTTCT TGTTTGGGTT	1260
	TOGAATGGAA ATATGGATAG CAGCATAAAG TACTTTTATT TTGACAAAAT TCATTTTTT	1320
55	CAACAATGGA GACATAGATT TGACCCACAA TAACTTCTCC CCCTCTCTTT TTACTCTGCT	1380
	CAAAAAGCAT CTCTCCTCCC ATTACCCAAC CTTGGTCATA AGTGTGCCTG GCTGGTTTGC	1440
60	AGATATTTCT TCTCCTTTCT AAAAATTCGC CATTAGTGCA TTTATTGAGA TGATCTCTAA	1500

	AGAGCTATGC CCTGACCTAC CCCTGATTCT ATGACATTGG GGCCCTTCTT TTGCTGAAAC	1560
5	TGCCTTACGT AATGGTTTTA CTCCTTGAAA GAGATTTGAC GGAATCCATT TTATGCCAAG	1620
5	TECTECCCTE CACTETTTCT GCAATATGTG GTGTATGCTG TEGTGATCTT GCTGGGAATG	1680
	ATTATAAGTG TGTGTGGT GGGGGAGTGG GTATTACATG CATTGCTGAA GAGTCAAAAA	1740
10	AAAAAAAAA AAACTCGA	1758
15	(2) INFORMATION FOR SEQ ID NO: 174:	
	(i) SEQUENCE CHARACTERISTICS:	
20	(A) LENGTH: 888 base pairs (B) TYPE: nucleic acid	:
20	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 174:	
25	CTGTTAGAAT GCCCAGTTTA CCTGGATGGC AACCCAACAG TGCTCCTGCC CACCTGCCCC	60
	TCAATCCTCC TAGAATTCAG CCCCCAATTG CCCAGTTACC AATAAAAACT TGTACACCAG	120
30	CCCCAGGGAC AGTCTCAAAT GCAAATCCAC AGAGTGASMC ACCACCTCGG GTAGAATTTG	180
50	ATGACAACAA TCCCTTTAGT GAAAGTTTTC AAGAACGGGA ACGTAAGGAA CGTTTACGAG	240
	AACAGCAAGA GAGACAACGG ATCCAACTCA TGCAGGAGGT AGATAGACAA AGAGCTTTGC	300
35	AGCAGAGGAT GGAAATGGAG CAGCATGGTA TGGTGGGCTC TGAGATAAGT AGTAGTAGGA	360
	CATCTGTGTC CCAGATTCCC TTCTACAGTT CCGACTTACC TTGTGATTTT ATGCAACCTC	420
10	TAGGACCCCT TCAGCAGTCT CCACAACACC AACAGCAAAT GGGGCAGGTT TTACAGCAGC	480
	AGAATATACA ACAAGGATCA ATTAATTCAC CCTCCACCCA AACTTTCATG CAGACTAATG	540
	AGCGAGGCAG GTAGGCCCTC CTTCATTTGT TCCTGATTCA CCATCAATCC CTGTTGGAAG	600
45	CCCAAATTT TCTTCTGTGA AGCAGGGACA TGGAAATCTT TCTGGGACCA GCTTCCAGCA	660
	GTCCCCAGTG AGGCCTTCTT TTACACCTGC TTTACCAGCA GCACCTCCAG TAGCTAATAG	720
50	CAGTCTCCCA TGTGGCCAAG ATTCTACTAT AACCCATGGA CACAGTTATC CGGGATCAAC	780
-	CCAATCGCTC ATTCAGTTGT ATTCTGATAT AATCCCAGAG GAAAAAGGGN AAAAAAAARA	840
	AMAARAAARA ARAAAGGAGA TGATGATGCA GAATTCCACC AAGGCTCC	888

(2) INFORMATION FOR SEQ ID NO: 175:

60 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2379 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 175:

	GGCAGAGCTA	GTGTGGACTC	CATCCCCCTG	GAGTGGGATC	ACGNCTATGA	CCTCAGTCGG	60
10	GACCTGGAGT	CTGCAATGTC	CAGAGCTCTG	CCCTCTGAGG	ATGAAGAAGG	TCAGGATGAC	120
	AAAGATTTCT	ACCTCCGGG	AGCTGTTGSC	TTATCAGGG	ACCACAGTGC	CCTAGAGTCA	. 180
15	CAGATCCGAC	AACTGGGCAA	AGCCTGGATG	ATAGCCGCTT	TCAGATACAG	CAAACCGAAA	240
15	ATATCATTCG	CAGCAAAACT	CCCACGGGGC	CGGAGCTAGA	CACCAGCTAC	AAAGGCTACA	300
	TGAAACTGCT	GGGCGAATGC	AGTAGCAGTA	TAGACTCCGT	GAAGAGACTG	GAGCACAAAC	360
20	TGAAGGAGGA	AGAGGAGAGC	CTTCCTGGCT	TTGTTAACCT	GCATAGTACC	GAAACCCAAA	420
	CGGCTGGTGT	GATTGACCGA	TGGGAGCTTC	TCCAGGCCCA	GGCATTGAGC	AAGGAGTTGA	480
25	GGATGAAGCA	GAACCTCCAG	AAGTGGCAGC	AGTTTAACTC	AGACTTGAAC	AGCATCTGGG	540
23	CCTGGCTGGG	GGACACGGAG	GAGGAGTTGG	AACAGCTCCA	GCGTCTGGAA	CTCAGCACTG	600
	ACATCCAGAC	CATCGAGCTC	CAGATCAAAA	AGCTCAAGGA	GCTCCAGAAA	GCTGTGGACC	660
30	ACCGCAAAGC	CATCATCCTC	TCCATCAATC	TCTGCAGCCC	TGAGTTCACC	CAGGCTGACA	720
	GCAAGGAGAG	CCGGGACCTG	CAGGATCGCT	TGTSGCAGAT	GAATGGGCGC	TGGGACCGAG	780
35	TGTGCTCTCT	GCTGGAGGAG	TGGCGGGGCC	TGCTGCAGGA	TGCCCTGATG	CAGTGCCAGG	840
33	GTTTCCATGA	AATGAGCCAT	GGTTTGCTTC	TTATGCTGGA	GAACATTGAC	AGAAGGAAAA	900
	ATGAAATTGT	CCCTATTGAT	TCTAACCTTG	ATGCAGAGAT	ACTTCAGGAC	CATCACAAAC	960
40	AGCTTATGCA	AATAAAGCAT	GAGCTGTTGG	AATCCCAACT	CAGAGTAGCC	TCTTTGCAAG	1020
	ACATGTCTTG	CCAACTACTG	GTGAATGCTG	AAGGAACAGA	CTGTTTAGAA	GCCAAAGAAA	1080
45	AAGTCCATGT	TATTGGAAAT	CGGCTCAAAC	TTCTCTTGAA	GGAGGTCAGT	CGTCATATCA	1140
43	AGGAACTGGA	GAAGTTATTA	GACGTGTCAA	GTAGTCAGCA	GGATTTGTCT	TCCTGGTCTT	1200
	CTGCTGATGA	ACTGGACACC	TCAGGGTCTG	TGAGTCCCAY	ATCAGGAAGG	AGCACCCCAA	1260
50	ACAGACAGAA	AACGCCACGA	GGCAAGTGTA	GTCTCTCACA	GCCTGGACCC	TCTGTCAGCA	1320
	GTCCACATAG	CAGGTCCACA	AAAGGTGGCT	CCGATTCCTC	CCTTTCTGAG	CCARGGCCAG	1380
55	GTCGGTCCGG	CCGCGGCTTC	CTGTTCAGAG	TCCTCCGAGC	AGCTCTTCCC	CTTCAGCTTC	1440
رر	TCCTGCTCCT	CCTCATCGGG	CTTGCCTGCC	TTGTACCAAT	GTCAGAGGAA	GACTACAGCT	1500
•	GTGCCCTCTC	CAACAACTTT	GCCCGGTCAT	TCCACCCCAT	GCTCAGATAC	ACGAATGGCC	1560
60	CTCCTCCACT	CTGAACTAAG	CAGATGCCAT	CTGCAGAAGT	GCTGGTAGCA	TAAGGAGGAT	1620

	CGGGTCATAA GCAATCCCAA ACTACCAACA AGAGGACCTT GATCTTGGCG AAAGCCMTCG	168
5	GTGTGGCAGC TTTAGCCTCC TCCAGATCAC ATGTGTGCAA ATTATGGCTT CAGAGGTGGA	174
J	AGATAAACAG TGACGGGGA ACAAACAGAC AACAAGAAGG TTTGGAAGAA ATCTGGTTTG	180
	AGACTCTGAA CCTTAGCACT AAGGAGATTG AGTAAGGACC TCCAAAGTTC CCCGGACTCA	186
10	TGAATTCTGG GCCCTTGGCC NATTCTGTGC ACAGCCAAGG ACTTCAGTAG ACCATCTGG	192
	CAGCTTTCCC ATGGTGCTGC TCCAACCATC AGATAAATGA CCCTCCCAAG CACCATGTCA	198
15	GTGTCGTACA ATCTACCAAC CAACCAGTGC TGAAGAGATT TTAGAACCTT GTAACATACA	204
15	ATTTTAAGA GCTTATATGG CAGCTTCCTT TTTACCTTGT TTTCCTTTGG GGCATGATGT	210
	TTTAACCTTT GCTTTAGAAG CACAAGCTGT AAATCTAAAA GGCACTTTTT TTTAGAGGTA	216
20	TAAAGAAAAA CTAGATGTAA TAAATAAGAT CATGGAAGGC TTTATGTGAA AAAAGTTGAA	222
	TGTTATAGTA AAAAAAAAG ATATTTATGT ATGTACAGTT TGCTAAAGCC AAGTTTTGTT	228
25	TGTATTGATT TCTTTGCATT TATTÄTAGAT ATTATAAAAT AAAAAAAAAA AAAAAAAAAC	234
-	TCGAGGGGG GCCCGGTACC CAATTCGCCC TATAGTGAG	237
30	(2) THEODMAINTON FOR CEO TO NO. 176.	
	(2) INFORMATION FOR SEQ ID NO: 176:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1348 base pairs	
35	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 176:	
40	GCGCCTTCAC GATGCCGGCG GTCAGTGGTC CAGGTCCCTT ATTCTGCCTT CTCCTCCTGC	6
	TCCTGGACCC CCACAGCCCT GAGACGGGGT GTCCTCCTCT ACGCAGGTTT GAGTACAAGC	12
45	TCAGCTTCAA AGGCCCAAGG CTGGCATTGC CTGGGGCTGG AATACCCTTC TGGAGCCATC	18
	ATGGAGGTGA GGGGCAGGGG TGGGGACCGC TATGCCCAGG GTCCCTCAAA GTGCTGGAGG	24
50	GGCTGTRACT TGGTGGGGAG TGGGTCTGTC ACAGCCATCC TCTGTCCAGG GTGGGGCAAG	30
JU		
	GCCTGGGACA GTGCCAGGCA CCCCAGGACC CCTTCCAGGC TTGTCTCCTG CTCCACCGCC	36
	GCCTGGGACA GTGCCAGGCA CCCCAGGACC CCTTCCAGGC TTGTCTCCTG CTCCACCGCC TCAACACCCC CCACCCCTGC CCAAGCTGTT TCTCCTCTGC CTCTCTNNTT CCCTGCCCCA	36 42
55	•	

	CTGGGGGCTA	CCTGGAGGGA	AGCATCCTCA	TCCCAGGTGA	GTGGGCACCA	GCCCTTCCCT	66
	GTATGTGTGT	TGTGGGTGGA	AGCAGGCATG	AGAGCATCTT	AGCCCATAGG	TTTGTATTCA	72
5	GGGACTTCCA	AACCCAGACC	TACAAAGAGT	GTGTCTTCTA	CCAGATCTTG	TTCAAAAAAG	780
	GGTTTGTGAT	GATGGAACTA	CACGATAGAG	GGAGTGAGCA	AGAACAATGA	GGATTAGAGT	840
10	GGAGCGTGAA	ATAGTCTAGG	AGCATGGCTT	CCAAAACATA	TGCTGTGAGG	TCTGTCCACC .	900
	TGAGAGTTGG	GCCATGGATT	TAATTCTGAG	CCTCTTAGCA	GGCAAAGCAA	AGACAGAAAG	960
	CAGATCGGCT	GTGGATTTCT	GTCTATAAAA	TCTGAGTTCT	TGGCCGGGTG	CGGTGGCTCA	1020
15	CGCCTGTAAT	CCCGCCCTT	TGGGAGGCCA	GGGCGGATGG	GTCGCGAGGT	CAGGAGGTTG	1080
	GAAACCATCC	TGGCCGGAAT	GGTGAAGCCC	TGACTCTACT	AGAAGTGCAA	AGATTGGCTG	1140
20	GGTGTGGTGG	CCTCCCCCTC	TGGTCCCAGC	TTCTCGGGAG	GCTGAGGCGG	GAGAGTTGCT	1200
20	TGGGCCTGGG	AGGCCGAGGT	TGCGGTGAGC	TGAGATCCTG	CCATTGCACT	TCAGCCTGGG	1260
	CACAGAGCCA	GACTCTGGCT	СААААААА	АААААААА	ACTCGAGGGG	GGCCCGTACC	1320
25	CAATTCGCCG	NATATGATCG	TAAACAAT				1348

## $30\,$ (2) information for SEQ ID No: 177:

35

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1502 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 177:

				•			
40	СТСААААТАА	ATAAATAAAT	AAAAATTTGT	ATTCCATTGA	TTTGGGTAGA	CACCAGGAAT	60
	GTGCATTTCT	AACAAGCTTT	CCAGGCGATC	CTATAGTAAG	TCATCTGTGG	ACTACTTTAA	120
45	GAAACTCTTC	TATAGAGAAT	GGAGTTGGAT	TAATAATAGG	TGATTTTTTA	CACTGGACTG	180
	ATTCACAAGA	ACCTAAACAG	TAGTCCATGA	AGCTGCTCAT	CTGTGGTAAC	TATTTGGCCC	240
	CGTCTCACTC	TGAAAGCAGC	AGGAGATGTT	GTTTACTTTG	TTTCTATCCC	CTTTGTCTGG	300
50	AGATTAATTT	TGGAATGAAA	GTTTTTCTCT	CTATGCCATT	CCTGGTTCTT	TTCCAAAGCC	360
	TCATACAAGA	GGATTAGGTC	ACAATGCATG	CATTACCTTT	TAAAAGAATG	CGATATTGAT	420
55	ACCGATGCTT	ACTITITITI	TTTTTNACTA	CTTGTTTTAT	TCCTTCCAGN	AAAGTATAGC	480
•	CCGCCTTTCT	ATAGCATAGT	TCTCTTTAGG	TGGAATGATT	CCTATAAGAT	TTCTCATTAT	540
	TAAATCATGC	ATTTTTCAAG	ATGGAATCAA	TMTTTGATTT	AATCTAAGCT	GATATTCTCA	600
60	TTTGTTAGAA	GAACAACCTA	CATGCTAGAG	AGAGAGGAGG	AAATATACCC	ACGACCACAC	660

	AGCCAGTTAG	TATCCAGTTG	GTGCTGGACT	CCAGCCAGGT	GTCCTGCCTC	ATGGTAGTTA	720
5	AATGATATAT	AGAAAAGGTA	AATTTTTAAA	GAAATATTTA	TTAATATATT	CCTATAAAAC	780
3	ATTTTAAAGG	TAACCACATA	AAAATGGTTA	ATTTTTCCAT	TCCAAAGTAA	ATGCTAAGCA	840
	TGTTTATTAA	TGAAGCAGTA	CTTCTGATTA	GTATATGACA	TTCTGAAGTT	AATTAAACTĆ	900
10	ATTGCACTAA	ATGTGTCTTC	CTTGGTATAG	TGGAGGATTT	GAGGATTGGA	ATATAGAGTA	960
	GAGTGCTTGC	TTAAGCCTGG	GAGCCCATCT	TTATAGCTAT	TTGATGTAAG	AAAAGAGACA	1020
15	TGGNCCATTT	СТАААСТАТА	TAAGGTGAGT	GTGTCTATTC	CCAGCAGATA	TAAAGGAAAA	1080
15	AGGAAACTTT	TTTGATTCCC	ACCTTCCCAG	CCTCACCTAG	CCATCTTCCA	GCCTCAAATA	1140
	TAGAGATGTT	AGTGCAAGGT	CCTGGGCTCT	AGGTGATCAT	TTCATAAGTC	CTTTACAGAT	1200
20	AAAGAAAAG	TAGTGTTTGT	ATGTTTGTTT	TTAAGTAACC	ССААААСААА	TTTATATTGT	1260
	ATTCAGCAAA	ATTGGAATTC	AGGTGTTTAA	TTTTAGAACA	TGAAGTGCCT	GCTGTTTTAA	1320
25	GCATTGACTT	GTATAAAAAG	AATTGCATGT	CTCCAGTAAG	CTTATGGGTT	TTCTCATTTT	1380
43	TAGGTATATG	GCTTTTAATC	ATGTAAAGTG	AAACATTAGT	TTTCTTGCAT	TTTATTACAG	1440
	GTTCTTTGTT	GCAATAAAGA	TGCTGCTGAA	ATTAATTGAA	ааааааааа	ÁAAAAAACTC	1500
30	GA						1502

#### 35 (2) INFORMATION FOR SEQ ID NO: 178:

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# (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1637 base pairs

- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 178:

45	ATTITCTAGC	CCACAAGGAC	TGAAGTTCAG	ATCCAAAAGT	TCACTTGCTA	ATTATCTTCA	60
	CAAAAATGGA	GAGACTTCTC	TTAAGCCAGA	AGATTTTGAT	TTTACTGTAC	TTTCTAAAAG	120
50	GGGTATCAAG	TCAAGATATA	AAGACTGCAG	CATGGCAGCC	CTGACATCCC	ATCTACAAAA	180
30	CCAAAGTAAC	AATTCAAACT	GGAACCTCAG	GACCCGAAGC	AAGTGCAAAA	AGGATGTGTT	240
	TATGCCGCCA	AGTAGTAGTT	CAGAGTTGCA	GGAGAGCAGA	GGACTCTCTA	ACTITACTIC	300
55	CACTCATTTG	CTTTTGAAAG	AAGATGAGGG	TGTTGATGAT	GTTAACTTCA	GAAAGGTTAG	360
	AAAGCCCAAA	GGAAAGGTGA	CTATTTTGAA	AGGAATCCCA	ATTAAGAAAA	CTAAAAAAGG	420
60	ATGTAGGAAG	AGCTGTTCAG	GTTTTGTTCM	AAGTGATAGC	AAAAGAGAAT	CTGTGTGTAA	480
UU							

	TAAAGCAGAT GCTGAAAGTG AACCTGTTGC ACAAAAAAGT CAGCTTGATA GAACTGTCTG	540
	CATTICTGAT GCTGGAGCAT GTGGTGAGAC CCTCAGTGTG ACCAGTGAAG AAAACAGCCT	600
5	TGTAAAAAAA AAAGAAAGAT CATTGAGTTC AGGATCAAAT TTTTGTTCTG AACAAAAAAC	660
	TTCTGGCATC ATAAACAAAT TTTGTTCAGC CAAAGACTCA GAACACAACG AGAAGTATGA	720
10	GGATACCTTT TTAGAATCTG AAGAAATCGG AACAAAAGTA GAAGTTGTGG AAAGGAAAGA	780
10	ACATTTGCAT ACTGACATTT TAAAACGTGG CTCTGAAATG GACAACAACT GCTCACCAAC	840
	CAGGAAAGAC TTCACTGAAG ATACCATCCC ACGGAACACA GATAGAAAGA AGGAAAACAA	900
15	GCCTGTATTT TTCCAGCAAA TATAACAAAG AAGCTCTTAG CCCCCACGA CGTAAAGCCT	960
•	TTAAGAAATG GACACCTCCT CGGTCACCTT TTAATCTCGT TCAAGAAACA CTTTTTCATG	1020
20	ATCCATGGAA GCTTCTCATC GCTACTATAT TTCTCAATCG GACCTCAGGC AAAATGGCAA	1080
20	TACCTGTGCT TTGGAAGTTT CTGGAGAAGT ATCCTTCAGC TGAGGTAGCA AGAACCGCAG	1140
	ACTGGAGAGA TGTGTCAGAA CTTCTTAAAC CTCTTGGTCT CTACGATCTT CGGGCAAAAA	1200
25	CCATTGTCAA GTTCTCAGAT GAATACCTGA CAAAGCAGTG GAAGTATCCA ATTGAGCTTC	1260
	ATGGGATTGG TGCACCCTGA AGACCACAAA TTAAATAAAT ATCATGACTG GCTTTGGGAA	1320
30	AATCATGAAA AATTAAGTCT ATCTTAAACT CTGCAGCTTT CAAGCTCATC TGTTATGCAT	1380
<b>J</b> 0	AGCTTTGCAC TTCAAAAAAG CTTAATTAAG TACAACCAAC CACCTTTCCA GCCATAGAGA	1440
	TTTTAATTAG CCCAACTAGA AGCCTAGTGT GTGTGCTTTC TTAATGTGTG TGCCAATGGT	1500
35	GGATCTTTGC TACTGAATGT GTTTGAACAT GTTTTGAGAT TTTTTTAAAA TAAATTATTA	1560
	ТТТСАСААСА АТССАААААА АААААААААА АААААААА	1620
40	AAAAAAAAA AAAAAAA	1637
	(2) INFORMATION FOR SEQ ID NO: 179:	
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2911 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
50	(D) TOPOLOGY: linear	•
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 179:	
55	GGTGGTTTTT GTTCTGCAAT AGGCGGCTTA GAGGGAGGGG CTTTTTCGCC TATACCTACT	60
	GTAGCTTCTC CACGTATGGA CCCTAAAGGC TACTGCTGCT ACTACGGGGC TAGACAGTTA	120
	CTGTCTCAGC TCTAGGATGT GCGTTCTTCC ACTAGAAGCT CTTCTGAGGG AGGTAATTAA	180
60	AAAACAGTGG AATGGAAAAA CAGTGCTGTA GTCATCCTGT AATATGCTCC TTGTCAACAA	240

	TGTATACATT CCTGCTAGGT GCCATATTCA TTGCTTTAAG CTCAAGTCGC ATCTTACTAG	300
_	TGAAGTATTC TGCCAATGAA GAAAACAAGT ATGATTATCT TCCAACTACT GTGAATGTGT	360
5	GCTCAGAACT GGTGAAGCTA GTTTTCTGTG TGCTTGTGTC ATTCTGTGTT ATAAAGAAAG	420
	ATCATCAAAG TAGAAATTTG AAATATGCTT CCTGGAAGGA ATTCTCTGAT TTCATGAAGT	. 480
0	GGTCCATTCC TGCCTTTCTT TATTTCCTGG ATAACTTGAT TGTCTTCTAT GTCCTGTCCT	540
	ATCTTCAACC AGCCATGGCT GTTATCTTCT CAAATTTTAG CATTATAACA ACAGCTCTTC	600
15	TATTCAGGAT AGTGCTGAAG ANGCGTCTAA ACTGGATCCA GTGGGCTTCC CTCCTGACTT	660
13	TATTITITGTC TATTGTGGCC TIGACTGCCG GGACTAAAAC TITACAGCAC AACTTGGCAG	720
	GACGTGGATT TCATCACGAT GCCTTTTTCA GCCCTTCCAA TTCCTGCCTT CTTTTCAGAA	780
20	ATGAGTGTCC CAGAAAAGAC AATTGTACAG CAAAGGAATG GACTTTTCCT GAAGCTAAAT	840
	GGAACACCAC AGCCAGAGTT TTCAGTCACA TCCGTCTTGG CATGGGGCAT GTTCTTATTA	900
25	TAGTCCAGTG TTTTATTTCT TCAATGGCTA ATATCTATAA TGAAAAGATA CTGAAGGAAG	960
23	GGAACCAGCT CACTGAARGC ATCTTCATAC AGAACAGCAA ACTCTATTTC TTTGGCATTC	1020
	TGTTTAATGG GCTGACTCTG GGCCTTCAGA GGAGTAACCG TGATCAGATT AAGAACTGTG	1080
30	GATTTTTTTA TGGCCACAGT GCATTTTCAG TAGCCCTTAT TTTTGTAACT GCATTCCAGG	1140
	GCCTTTCAGT GGCTTTCATT CTGAAGTTCC TGGATAACAT GTTCCATGTC TTGATGGCCC	1200
35	AGGTTACCAC TGTCATTATC ACAACAGTGT CTGTCCTGGT CTTTGACTTC AGGCCCTCCC	1260
<i>33</i>	TGGAATTTTT CTTGGAAGCC CCATCAGTCC TTCTCTCTAT ATTTATTTAT AATGCCAGCA	1320
	AGCCTCAAGT TCCGGAATAC GCACCTAGGC AAGAAAGGAT CCGAGATCTA AGTGGCAATC	1380
40	TTTGGGAGCG TTCCAGTGGG GATGGAGAAG AACTAGAAAG ACTTACCAAA CCCAAGAGTG	1440
	ATGAGTCAGA TGAAGATACT TTCTAACTGG TACCCACATA GTTTGCAGCT CTCTTGAACC	1500
15	TTATTTTCAC ATTTTCAGTG TTTGTAATAT TTATCTTTTC ACTTTGATAA ACCAGAAATG	1560
45	TTTCTAAATC CTAATATTCT TTGCATATAT CTAGCTACTC CCTAAATGGT TCCATCCAAG	162
	GCTTAGAGTA CCCAAAGGCT AAGAAATTCT AAAGAACTGA TACAGGAGTA ACAATATGAA	168
50	GAATTCATTA ATATCTCAGT ACTTGATAAA TCAGAAAGTT ATATGTGCAG ATTATTTTCC	174
	TTGGCCTTCA AGCTTCCAAA AAACTTGTAA TAATCATGTT AGCTATAGCT TGTATATACA	180
56	CATAGAGATC AATTTGCCAA ATATTCACAA TCATGTAGTT CTAGTTTACA TGCCAAAGTC	186
55	TTCCCTTTTT AACATTATAA AAGCTAGGTT GTCTCTTGAA TTTTGAGGCC CTAGAGATAG	192
	TCATTTTGCA AGTAAAGAGC AACGGGACCC TTTCTAAAAA CGTTGGTTGA AGGACCTAAA	198
60	TACCTGGCCA TACCATAGAT TTGGGATGAT GTAGTCTGTG CTAAATATTT TGCTGAAGAA	204

	GCAGTTTCTC	AGACACAACA	TCTCAGAATT	TTAATTITTA	GAAATTCATG	GGAAATTGGA	2100
5	TTTTTGTAAT	AATCTTTTGA	TGTTTTAAAC	ATTGGTTCCC	TAGTCACCAT	AGTTACCACT	2160
3	TGTATTTTAA	GTCATTTAAA	CAAGCCACGG	TGGGGCTTTT	TTCTCCTCAG	TTTGAGGAGA	2220
	AAAATCTTGA	TGTCATTACT	CCTGAATTAT	TACATTTTGG	AGAATAAGAG	GGCATTTTAT	2280
0.	TTTATTAGTT	ACTAATTCAA	GCTGTGACTA	TTGTATATCT	TTCCAAGAGT	TGAAATGCTG	2340
-	GCTTCAGAAT	CATACCAGAT	TGTCAGTGAA	GCTGATGCCT	AGGAACTTTT	AAAGGGATCC	2400
<b>c</b>	TTTCAAAAGG	ATCACTTAGC	AAACACATGT	TGACTTTTAA	CTGATGTATG	AATATTAATA	2460
.5	СТСТАААААТ	AGAAAGACCA	GTAATATATA	AGTCACTTTA	CAGTGCTACT	TCACACTTAA	2520
	AAGTGCATGG	TATTTTTCAT	GGTATTTTGC	ATGCAGCCAG	TTAACTCTCG	TAGATAGAGA	2580
20	AGTCAGGTGA	TAGATGATAT	TAAAAATTAG	CAAACAAAAG	TGACTTGCTC	AGGGTCATGC	2640
	AGCTGGGTGA	TGATAGAAGA	GTGGGCTTTA	ACTGGCAGGC	CTGTATGTTT	ACAGACTACC	2700
25	ATACTGTAAA	TATGAGCTTT	ATGGTGTCAT	TCTCAGAAAC	TTATACATTT	CTGCTCTCCT	2760
2.5	TTCTCCTAAG	TTTCATGCAG	ATGAATATAA	GGTAATATAC	TATTATATAA	TTCATTTGTG	2820
	ATATCCACAA	TAATATGACT	GGCAAGAATT	GGTGGAAATT	TGTAATTAAA	ATAATTATTA	2880
30	AACCTAAAAA	AAAAAAAA.	AAAAACTCGA	G			291

## 35 (2) INFORMATION FOR SEQ ID NO: 180:

40

#### (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 519 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
  - (D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 180:

	•						
45	GGCACGAGCC	CCAGGCCAGC	CAGGGCCAGG	CCTACTTTGG	CCACCCTTAA	ATTAGAATGT	60
	GGGGTCAGGG	GTCACAGAAA	AGCCATTTCT	CTGACCTAGT	GTTTGGCGTC	CGGGAACTCT	120
50	GTGCCCAACC	TTCAGACCCT	GGCAGTCCTC	ACTGAGGCCA	TTGGCCCAGA	GCCCGCCATC	180
30	CCCCGARACC	CCCGGGAGCC	GCCTGTTGCC	ACGTCCACAC	CTGCCACACC	CTCTGCCGGG	240
	CCCCAGCCCC	TCCCAACCGG	GACCGTGCTG	GTCCCTGGGG	GICCIGCCC	ACCTTGCCTT	300
55	GGGGAGGCAT	GGGCCCTCCT	CCTCCCACCC	TGCCGGCCGT	CACTCACCTC	TTGCTTCTGG	360
	TCCCCCAGGC	CTAGCCCTTG	GAAGGAGACA	GGAGTCTAGG	GAGGCTGAAG	CCCACTCCCG	420
40	GGGAGGCCCG	TGCTCCTCCA	GCCCCAGGGA	CAGCAAGGAA	AAGAGAAGAG	AGCAGAGCAT	480

# TTCATGGCTC TAATAAAAAA AAAAAAAAAA AAAACTCGA

519

10

(2) INFORMATION FOR SEQ ID NO: 181:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 968 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 181:

	•		•			•	
15	TCCCCTTGGG	GCCGGAAAAA	GCGGGGTTGG	CCTGNCCATT	GGTTNTCCAT	GCCGCCCGCC	60
	CATGCCCCAG	TACTAGCCTG	CAGTCCCAAT	GTAGCCCCTC	CCTCYTCCMA	GAGCCCYTCM	120
20	AACCGCCCCG	STCANTIGTG	ATTTCAGGAG	GATTTGATGA	AGATGTTAAA	GCGAAAGTGG	180
	AGAACCTTCT	CGGGATTTCC	AGCCTGGAAA	AAACGGACCC	TGTTAGGCAA	GCACCCTGCA	240
25	GCCCTCCCTG	TCCCCTTCTT	CCCCTCCCCT	TCYCCCGCCC	GTGGAGACAG	CTGTTYTCAG	300
23	CAGGGCTCTC	CGCAGGGAGG	GGGCCGGCTC	CTTCCCTGGC	AGCAACATCC	TIGCCCTIGT	360
	CACACAAGTC	AGCCTCCATC	TGCGCAGCTC	TGTGGATGCG	CTGCTGGAGG	GCAACAGGTA	420
30	TGTCACTGGC	TGGTTCAGCC	CCTACCACCG	CCAGCGGAAG	CTCATCCACC	CGGTCATGGT	480
	TCAGCACATC	CAGCCCGCAG	CGCTCAGCCT	CCTGGCACAG	TGGAGCACCC	TCGTGCAGGA	540
35	GCTGGAGGCT	GCCCTGCAGC	TGGCTTTCTA	CCCGGATGCC	GTGGAGGAGT	GGCTGGAGGA	600
33	AAACGTGCAC	CCCAGCCTGC	AGCGGCTGCA	ARCTCTGCTG	CAGGACCTCA	GCGAGGTGTC	660
	TGCCCCCCC	CTGCCACCCA	CCACCCCTGG	CAGGGACGTT	GCTCAGGACC	CCTGAGGGGA	720
40	GAGCTCATGC	CAGGGGGCTC	CTGCTGGAGG	CTGGGGGGG	TCTGCWYTKY	CWWWIGGCCT	780
	GGGCAATACG	GCCCACGTGG	GCGTCGTGCC	: CTCTGGCCCA	GCAGTGTCTT	GCCCACACTC	840
45	AGTTCCTGAG	GCCCTGGGC	: AGCCCCTGGG	GGAGAGACTA	GAAAACACAG	AAGGAAGCAG	900
40	CACAGGGAGA	CCCCCTTTGI	GATCTGCATC	TGTGACACTG	ATTCTTTGGA	AATAAAGAGT	960
	GGAAGCTG						968

50

#### (2) INFORMATION FOR SEQ ID NO: 182:

55 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1128 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(XI) SEQUENCE DESCRIPTION. SEQ 15 No. 101.					
•	TGTAAAAGTT ATCAGTAATC CTAATTCTTT TCCTGGGTTT TCCTTTTGTC ACTTATTAAT	60				
5	CAGTTTTTGA AAGGACGAAT GAATTTAGAG ATGTACTCTG GAGCAGTATC ATGTTAAACC	120				
	AGGGGTATAT TAGAAAAATC ATCCTCATAA TCATTCTGGG AAGTTTTTCC TCCCCAAAAA	180				
••	AAGCCATCCT GATGGGTTTT CAAAACCAGA AAAAAGCTCT TAATGAGGAA CAGACCACTG	240				
10	GAGTACCCAT GAGCATCTCA GGAAAACTGA GACCCTCGAG AAGCCTTGAT TTCGTGCAAC	300				
	CCCCAAGGTT TCAGAGCCAG CAGCCCAGTG CTGTGGTTGA CAGACGTGGT TTTKTGGRGA	360				
15	AAGCAGCCAG AGGCCAGGAA TTTTCAGAGT CGTGAGTCAC GRTYTCCCAC CCAAGATTAG	420				
	AGCAMAGATT AGCCATACTG AGATTTGGTA AAATCATTCT GTCTAAGCAA TGGAGGTGTG	480				
20	TGCAMACGTG CAGTGCCTGT TCACAGGGGA TGCAGGCAGA TCSYGGGTTT AGGATGGGGR	540				
20	AGGCCACCGC ACCCCCYTTC AYTGCTCTGC ACCTGCTCCC TCACGTGGAC ACTGTCCACA	600				
	ACTGTGGCTC TCACAGGACA GTTGCCCAAG GAGCTCATAT CTTATTGGAG ATAGGGGGTC	660				
25	GTACAGGTGA CATTCATGAG CAGTGTGAGC CGGGTGACAT GGGGGTGTCA ACCCAGCATC	720				
	TGTCCAGGAG CTCCTCCTGC AGCGGCTCTG GCAGGTGGCC TGAGGCTCCT TTTTGAGAGA	780				
30	GAACTGTTTG GCCTTCCTGT CTCCTCTCCT CTGATCTGTT CTTTCTTGGA ACACCACCCA	840				
30	AGAACGTCAC CTCCTCCATC AGATTGTGAG CTCCTGGAGG GCAGGAGCTG TGTCCTTCTA	900				
	TTCATCTTCC TATCCCCAGA ACCTTGCACA GATCCTGGAA TGTGGTAGGT GCTCAGTAAA	960				
35	TGTGTGTTGA ATAAATGAAT GAATGAATGA ACAAATGAAT GAATTTGCTT ACTTCAAGGC	1020				
	AAAAGAACCA TGAAACTGTA TTTTGAGTTT CTATGTTATA GCAGTCAGCA AATCCTATTA	1080				
40	AATACTTTGT GTTTCCAAGC AAAAAAAAAA AAAAAAAAA AAACTCGA	1128				
40						
	(2) INFORMATION FOR SEQ ID NO: 183:					
45	(i) SEQUENCE CHARACTERISTICS:					
	(A) LENGTH: 2276 base pairs (B) TYPE: nucleic acid					
	(C) STRANDEDNESS: double					
50	(D) TOPOLOGY: linear					
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 183:					
55	CCGCGGCGTC TGACCTCATG GCGTAGAGCC TAGCAACAGC GCAGGCTCCC AGCCGAGTCC	60				
33	GTTATGGCCG CTGCCGTCCC GAAGAGGATG AGGGGGCCAG CACAAGCGAA ACTGCTGCCC	120				
	GGGTCGGCCA TCCAAGCCCT TGTGGGGTTG GCGCGGCCGC TGGTCTTGGC GCTCCTGCTT	180				
60	GTGTCCGCCG CTCTATCCAG TGTTGTATCA CGGACTGATT CACCGAGCCC AACCGTACTC	240				

	AACTCACATA TTTCTACCCC AAATGTGAAT GCTTTAACAC ATGAAAACCA AACCAAACCT	300
c	TCTATTTCCC AAATCAGCAC CACCCTCCCT CCCACGACGA GTACCAAGAA AAGTGGAGGA	360
5	GCATCTGTGG TCCCTCATCC CTCGCCTACT CCTCTGTCTC AAGAGGAAGC TGATAACAAT	420
	GAAGATCCTA GTATAGAGGA GGAGGATCTT CTCATGCTGA ACAGTTCTCC ATCCACAGCC	480
10	AAAGACACTC TAGACAATGG CGATTATGGA GAACCAGACT ATGACTGGAC CACGGGCCCC	540
	AGGGACGACG ACGAGTCTGA TGACACCTTG GAAGAAAACA GGGGTTACAT GGAAATTGAA	600
٠,٠	CAGTCAGTGA AATCTTTTAA GATGCCATCC TCAAATATAG AAGAGGAAGA CAGCCATTTC	660
15	TTTTTCATC TTATTATTTT TGCTTTTTGC ATTGCTGTTG TTTACATTAC ATATCACAAC	720
	AAAAGGAAGA TTTTTCTTCT GGTTCAAAGC AGGAAATGGC GTGATGGCCT TTGTTCCAAA	780
20	ACAGTGGAAT ACCATCGCCT AGATCAGAAT GTTAATGAGG CAATGCCTTC TTTGAAGATT	840
	ACCAATGATT ATATTTTTTA AAGCACTGTG ATTTGAATTT GCTTATGTAA TTTTATTTGC	900
25	TTGACTTTT ATATGATATT GTGCAAATGT TTGCCATAGG CAATTGGTAC TTAAATGAGA	960
25	GGTGAGTCTC TCTTTTGCCT TGGTGCTTTG GAAATTAAAT GTCACAAACG AGTATATAAT	1020
	TTTTTATCTG TACTTTTAGA GCTGAGTTTA ATCAGGTGTC CAAAATGTGA GTTAAACATT	1080
30	ACCTTATATT TACACTGTTA GTTTTTATTG TTTTAGATTT ATTATGCTTC TTCTGGAAGT	1140
	ATTAGTGATG CTACTTTTAA AAGATCCCAA ACTTGTAACT AAATTCTGAC ATATCTGTTA	1200
35	CTGCTGACTC ACATTCATTC TCCGCCATTC AAATACTATT TTTTATCCAC ATTTTTTTTT	1260
33	GTTCCCAAAC TGTAATGTAC AAGGATATGT GTGATAATGC TTTGGATTTG AGTAATATTT	1320
	TTTTTTCTTC CAAGAAAACT GCTTTGGATA TTTTTAGATA ATTTAAACAT AATTTAGGAT	1380
40	AATGATATTG CTCAATCTGA CCACAATTTT AGGTAAAACA TTAAATGTGT CAGAAATCTT	1440
	GGCAACAGAG ACTCTGCAGC TTGCAGTGGA CATAGATAAA ATGTTACAGA GATACTATTT	1500
45	TTTTGGTTGG AATTACTATA TTAAATTTAG AAGCAGAAAC TGGTAAAATG TTAAATACAT	1560
43	GTACAATTGC TTTTAGTTAG CAATTGATTG TAGCATGGGT TCCTCCAAGG TTTCAAGCAA	1620
	TGGGCAGAGT TTAAAATTAT ATCAGATTCG TTTACTTCGT TTATTATTTT ACAGTAAATT	1680
50	TGAATAAATC TTAGGGGTCA TTATCACTTA AATAATACTG TACCTAGGTC TTTCAAATTA	1740
	AAATTATACC TGAATGAAGT TGTTTGTATA CATAAAGGAT ATTTGTGTAC AATTACCTTT	1800
55	TTTCCCCCAC ACTIGITITC TTTGTTTTTG TTTTTTATGG CAACTGGAAA GTATTTACTA	1860
JJ	TGGGATTCAT TTATGTCTGT CTTTCTATCA TAAAGAATTG ATCAATATGT AAATATGTGA	192
	TTTGAACCAT GGTTGACTTA CAAGTGTCAC TACAGCTTTT TAGAAAACAT AGCCCTAATA	198
60	TATGTTAAGC AGGACCCGGG TGAGCCAGTG GGCTTGCGCT TTATGTAGAG CTGGAAGAAG	204

	GCCGTCCATC CTGTCTCTTG GGCGGACAGT GTACTTTCCT AATAGGGAAG GGAAGCACAA	2100
		2160
_	TGGAAATACC CCTGAACCGT TTTATTGCAG TAATTTTTTT CATATCTGAA ACTATTATTT	2160
5	AATATTTTGA ATAAGATTTT AAAAAATAAA TGGCAAAGAT ATAAATCTAA AAAAAAAAAA	2220
	АААААА ААААААААА ААААААААА ААААААААА АААА	2276
10		
	(2) INFORMATION FOR SEQ ID NO: 184:	•
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2500 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double	
00	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 184:	
	TCCAAGCTAC GCCACTCGGG CTGGGGCGTT GGGAGCGGGA GTGCAGAGCG TGGTCGTGGC	60
25	GGCGCGGTG AGAAGAGCGA GGCGKAGGAG GGGGTGCCAT GGCCGGGCAG CAGTTCCAGT	120
	ACGATGACAG TGGGAACACC TTCTTCTACT TCCTCACCTC CTTCGTGGGG CTCATCGTGA	180
	TCCCGGCGAC ATACTACCTC TGGCCCCGAG ATCAGAATGC CGAGCAAATT CGATTAAAGA	240
30	ATATCAGAAA AGTATATGGA AGGTGTATGT GGTACGTTTA CGGTTATTAA AACCCCAGCC	300
	AAATATTATT CCTACAGTAA AGAAAATAGT TCTGCTTGCA GGATGGGCAT TGTTCTTATT	360
35	CCTTGCATAT AAAGTTTCCA AAACAGACCG AGAATACCAA GAATACAATC CTTATGAAGT	420
	ATTAAATTTG GATCCTGGAG CCACAGTAGC AGAAATTAAA AAACAATATC GTTTGCTGTC	480
	ACTTAAATAT CATCCAGATA AAGGAGGTGA TGAGGTTATG TTCATGAGGA TAGCAAAAGC	540
40	TTATGCTGCT TTAACGGATG AAGAGTCCCG GAAAAATTGG GAAGAATTTG GAAATCCAGA	600
	TGGGCCTCAA GCCACAAGCT TTGGAATTGC CCTGCCAGCT TGGATAGTTG ACCAGAAAAA	660
45		720
	GGGCTCTTGG TGGTATCGCT CAATACGCTA TAGTGGAGAC CAGATTCTAA TACGSACAAC	780
	ACAGATTTAT ACATACTTTG TTTATAAAAC CCGAAATATG GATATGAAAC GTCTTATCAT	
50	GGTTTTGGST GGAGCTTCTG AATTTGATCC TCAGTATAAT AAAGATGCCA CAAGGATGC	900
•	AACGGATAAT ATTCTAATAC CACAGCTAAT CAGAGAAATT GGCAGCATTA ATTTAAAGAA	960
5:	5 GAATGAGCCT CCACTTACCT GCCCATATAG CCTGAAGGCC AGAGTTCTTT TACTGTCTCA	1020
	TCTTGCTAGA ATGAAAATTC CTGAGACCCT TGAAGAAGAT CAGCAATTCA TGCTAAAAAA	
_	GTGTCCTGCC CTACTTCAAG AAATGGTTAA TGTAATCTGC CAACTAATAG TAATGGCCCG	1140

	GAACCGTGAA G	BAAAGGGAGT '	TTCGTGCTCC	AACTTTGGCA	TCCCTAGAAA	ACTGCATGAA	1200
	GCTTTCTCAG A	ATGGCCGTTC	AGGGACTTCA	GCAATTTAAG	TCTCCCCTTC	TGCAGCTCCC	1260
5	TCATATTGAA (	GAGGACAATC	TTAGACGGGT	TTCTAATCAT	AAGAAGTATA	AAATTAAAAC	1320
	TATCCAGGAT T	ITGGTGAGTT	TAAAAGAATC	AGATCGTCAC	ACTCTACTGC	ACTTCCTTGA	1380
10	AGATGAAAAA 1	TATGAAGAGG	TTATGGCTGT	CCTTGGGAGT	TTTCCATATG	TGACCATGGA .	1440
10	TATAAAATCA (	CAGGTGTTAG	ATGATGAAGA	TAGCAACAAC	ATCACAGTAG	GATCCTTAGT	1500
	TACAGTGTTG (	GITAAGTIGA	CAAGGCAAAC	AATGGCTGAA	GTATTTGAAA	AGGAGCAGTC	1560
15	CATCTGTGCT	GCAGAGGAAC	AGCCAGCAGA	AGATGGGCAG	GGTGAAACTA	ACAAGAACAG	1620
	GACAAAAGGA (	GGATGGCAAC	AGAAGAGTAA	AGGACCCAAG	AAAACTGCTA	AATCAAAAAA	1680
20	AAAGAAACCT '	TTAAAAAAAA	AACCTACACC	TGTGCTATTA	CCACAGTCAA	AGCAACAGAA	1740
20	ACAAAAGCAG	GCAAATGGAG	TCGTTGGGAA	TGAAGCTGCA	GTAAAGGAAG	ATGAAGAAGA	1800
	AGTITCAGAT	AAGGGCAGTG	ATTCTGAAGA	AGAAGAAACC	AATAGAGATT	CCCAAAGTGA	1860
25	GAAAGATGAT	GGTAGTGACA	GAGACTCTGA	TAGAGAGCAA	GATGAAAAAC	AAAACAAAGA	1920
•	TGATGAAGCA	GAGTGGCAAG	AATTACAACA	AAGCATACAG	CGAAAAGAGA	GAGCICTATT	1980
30	GGAAACCAAA	TCAAAAATAA	CACATCCTGT	GTATAGCCTT	TACTTTCCTG	AGGAAAAACA	2040
30	AGAATGGTGG	TGGCTTTACA	TTGCAGATAG	GAAGGAGCAG	ACATTAATAT	CCATGCCATA	2100
	TCATGTGTGT	ACGCTGAAAG	ATACAGAGGA	GGTAGAGCTG	AAGTTTCCTG	CACCAGGCAA	2160
35	GCCTGGAAAT	TATCAGTATA	CIGIGITICI	GAGATCAGAC	TCCTATATGG	GTTTGGATCA	2220
	GATTAAACCA	TTGGAAGTTK	GGAAGTTCAT	GAGGCTGAAG	CCTGTGCCAG	; AAAATCACCC	2280
40	ACAGTGGGAT	ACAGCAATAG	AGGGGGATGA	AGACCAGGAG	GACAGTGAGG	GCTTTGAAGA	2340
→∪	TAGCTTTGAG	GGAGGAAGAG	GGAGGGAGGA	AGGAAGGTGG	TGGACTTAAG	GCAGTTACTC	2400
	TGGAATGGGA	CCCACAGTGT	TTTGCACCAT	ATTTTGGCAP	TTTTTTTGG	CCGTTTTTNG	246
45	GAAGTGTTTT	CCNTNAANCC	CAGGAACCAT	TACAGAACCO	<b>;</b>		250

## 50 (2) INFORMATION FOR SEQ ID NO: 185:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1337 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 185:

60 CTTCCGGTTC TCCGGGCAGC TGCCACTGCT GTAGCTTCTG CCACCTGCCA CGACCGGGCC

•	TCTCCCTGGC GTTTGGTCAC CTCTGCTTCA TTCTCCACCG CGCCTATGGT CCCTCTTGGA	120
-	GCCAGCGTGG CGGGCCTGGC GGCTCCCGGG TGGTGAGAGA GCGGTCCGGG AACGATGAAG	180
5	GCCTCGCAGT GCTGCTGCTG TCTCAGCCAC CTCTTGGCTT CCGTCCTCCT CCTGCTGTTG	240
	CTGCCTGAAC TAAGCGGGYC CCTGGMAGTC CTGCTGCAGG CAGCCGAGGC CGCGCCAGGT	300
10	CTTGGGCCTC CTGACCCTAG ACCACGGACA TTACCGCCGC TGCCACCGGG CCCTACCCCT	360
	GCCCAGCAGC CGGGCCGTGG ȚCTGGCTGAA GCTGCGGGGC ÇGCGGGGCTC CGAGGGAGGC	420
15	AATGGCAGCA ACCCTGTGGC CGGGCTTGAG ACGGACGATC ACGGAGGGAA GGCCGGGGAA	480
13	GCCTCGGTGG GTGGCGGCCT TGCTGTGAGC CCCAACCCTG GCGACAAGCC CATGACCCAG	540
	CCGCCCTGA CCGTGTTGAT GGTGGTGAGC GGCGCGGTGC TGGTGTACTT CGTGGTCAGG	600
20	ACGGTCAGGA TGAGAAGAAG AAACCGAAAG ACTAGGAGAT ATGGAGTTTT GGACACTAAC	660
	ATAGAAAATA TOGAATTGAC ACCTTTAGAA CAGGATGATG AGGATGATGA CAACACGTTG	720
25	TTTGATGCCA ATCATCCTCG AAGATAAGAA TGTGCCTTTT GATGAAAGAA CTTTATCTTT	780
23	CTACAATGAA GAGTGGAATT TCTATGTTTA AGGAATAAGA AGCCACTATA TCAATGTTGG	840
	GGGGGTATTT AAGTTACATA TATTTTAACA ACCTTTAATT TGCTGTTGCA ATAAATACCG	900
30	TATCCTTTTA TTATATCTTT ATATGTATAG AAGTACTCTR TTAATGGGCT CAGAGATGTT	960
	GGGGATAAAG TATACTGTAA TAATTTATCT GTTTGAAAAT TACTATAAAA CGGTGTTTTC	1020
35	TGATCGGTTT TTGTTTCCTG CTTACCATAT GATTGTAAAT TGTTTTATGT ATTAATCAGT	1080
33	TAATGCTAAT TATTTTTGCT GATGTCATAT GTTAAAGAGC TATAAATTCC AACAACCAAC	1140
	TGGTGTGTAA AAATAATTTA AAATTTCCTT TACTGAAAGG TATTTCCCAT TTTTGTGGGG	1200
40	AAAAGAAGCC AAATTTATTA CTTTGTGTTG GGGTTTTTAA AATATTAAGA AATGTCTAAG	1260
-	TTATTGTTTG CAAAACAATA AATATGATTT TAAATTCTCT TAAAAAAAAA AAAAAAAACC	1320
45	CCGGGGGGGG GCCCGGN	1337

(2) INFORMATION FOR SEQ ID NO: 186:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 941 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 186:

GCCACGAGCC TGGACGCAGC AGCCACCGCC GCGTCCCTCT CTCCACGAGG CTGCCGGCTT

	AGGACCCCCA	GCTCCGACAT	GTCGCCCTCT	GGTCGCCTGT	GTCTTCTCAC	CATCGTTGGC	120
	CTGATTCTCC	CCACCAGAGG	ACAGACGTTG	AAAGATACCA	CGTCCAGTTC	TTCAGCAGAC	180
5	TCAACTATCA	TGGACATTCA	GGTCCCGACA	CGAGCCCCAG	ATGCAGTCTA	CACAGAACTC	240
	CAGCCCACCT	CTCCAACCCC	AACCTGGCCT	GCTGATGAAA	CACCACAACC	CCAGACCCAG	300
10	ACCCAGCAAC	TGGAAGGAAC	GGATGGGCCT	CTAGTGACAG	ATCCAGAGAC	ACACAAGAGC	360
10	ACCAAAGCAG	CTCATCCCAC	TGATGACACC	ACGACGCTCT	CTGAGAGACC	ATCCCCAAGC	420
	ACAGACGTCC	AGACAGACCC	CCAGACCCTC	AAGCCATCTG	GTTTTCATGA	GGATGACCCC	480
15	TTCTTCTATG	ATGAACACAC	CCTCCGGAAA	CGGGGGCTGT	TGGTCGCAGC	TGTGCTGTTC	540
	ATCACAGGCA	TCATCATCCT	CACCAGTGGC	AAGTGCAGGC	AGCTGTCCCG	GTTATGCCGG	600
20	AATCATTGCA	GGTGAGTCCA	TCAGAAACAG	GAGCTGACAA	CCYGCTGGGC	ACCCGAAGAC	660
20	CAAGCCCCCT	GCCAGCTCAC	CGTGCCCAGC	CTCCTGCATC	CCCTCGAAGA	GCCTGGCCAG	720
	AGAGGGAAGA	CACAGATGAT	GAAGCTGGAG	CCAGGGCTGC	CGGTCCGAGT	CTCCTACCTC	780
25	CCCCAACCCT	GCCCGCCCCT	GAAGGCTACC	TGGCGCCTTG	GGGGCTGTCC	CTCAAGTTAT	840
	CTCCTCTGYT	AAGACAAAAA	GTAAAGCACT	GTGGTCTTTG	СААААААААА	AAAAAAAA	900
30	AAAAAAAAA	AAAAAAAAA.	AAAAAAAA.	AAAAAACTCG	A	·.	943

(2) INFORMATION FOR SEQ ID NO: 187:

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#### (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 654 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- 40 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 187:

45	GAATTCGGCA	CGAGGCAGCT	TGTGCTTTAA	AGGAGGTGTT	CAAAGCATGT	CTGAGCAGAG	60
43	ACTITIGGGC	TCTGTTTTAA	TTAATACTTT	AAAATAATTC	ATATTTAAAA	TATCARATGT	120
	TTCCATAAAG	AGGAGGATGT	TTAAATGCCT	CCAGACTACA	TTCCTTTTTA	TTSCTTGATT	180
50	TTACCTGGGA	GTCCAAAGTT	CAATTCCCAT	AAAGCAAGCG	TTTTATTIGT	CACTTTCAAT	240
	ATACATCCGA	TTGCCATGCT	TAAGATGCAA	TATGGGCTGC	GGAAATAGGT	TAACCCACAG	300
55	GCTCCCAGGG	CCCAGTGTAG	AAGGTGAGAG	ATTCGTGTAA	AATGATTCAA	ATAAAAGGAA	360
<b>33</b>	GACCCTGGCC	GGGTGCCGTA	RCTCACGCCT	GTAATCCCAG	CACTTTGGGA	GGCCGAAGCG	420
	AGTGGATGAC	GAGGTTAGGA	GTTGGAGACC	AGCCTGGCCA	ACATCGTGAA	ACCCCCTCTC	480
60	ТАСТАААААТ	ACAAAAATTA	GCCGGGCATG	GTGGCAGGCA	CCTGTAATCC	TAGCTAGTTG	540

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	GGAGGCTGAG GCAGGAGAAT CGTTTGAATC TGGGAGTTGG AGGTTGTCAG TGAGCTGAGA	600
5	TCGCGCCACA GCACTCCAGC CTGGGTGACA GGGTGAGACT CTGTCTCAAA NAGA	654
10	(2) INFORMATION FOR SEQ ID NO: 188:  (i) SEQUENCE CHARACTERISTICS:	-
15	(A) LENGTH: 1848 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 188:	
20	GAAACTGGAC CGGAGAACCG GAGCGAAGCG AAGCGGAAGC CCGGAATGAG GCCGGACTGG	60
20	AAAGCCGGAG CGGGCCAGG CGGGCCTCCC CAAAAGCCTG CCCCTTCATC CCAGCGGAAA	120
	CCGCCGGCCC GGCCGAGCGC GGCGGCCGCT GCGATTGCAG TCGCGGCGGC GGAGGAAGAG	180
25	AGACGGCTCC GGCAGCGGAA CCGCCTGAGG CTGGAGGAGG ACAAACCGGC CGTGGAGCGG	240
	TECTTGGAGG AGCTGGTCTT CGGCGACGTC GAGAACGACG AGGACGCGTT GCTGCGGCGT	300
30	CTGCGAGGCC CGAGGGTTCA AGAACATGAA GACTCGGGTG ACTCAGAAGT GGAGAATGAA	360
30	GCAAAAGGTA ATTTTCCACC TCAAAAGAAG CCAGTTTGGG TGGATGAAGA AGATGAAGAT	420
	GAGGAAATGG TTGACATGAT GAACAATCGG TTTCGGAAGG ATATGATGAA AAATGCTAGT	480
35	GAAAGTAAAC TITCGAAAGA CAACCTTAAA AAGAGACTTA AAGAAGAATT CCAACATGCC	540
	ATGGGAGGAG TACCTGCCTG GGCAGAGACT ACTAAGCGGA AAACATCTTC AGATGATGAA	600
40	AGTGAAGAGG ATGAAGATGA TTTGTTGCAA AGGACTGGGA ATTTCATATC CACATCAACT	660
40	TCTCTTCCAA GAGGCATCTT GAAGATGAAG AACTGCCAGC ATGCGAATGC TGAACGTCCT	720
	ACTGTTGCTC GGATCTCCAT CTGTGCAGTT CCATCCCGGT GCACAGATTG TGATGGTTGC	780
45	TGGGATTAGA TAATGCTGTA TCACTATTTC AGGTTGATGG GAAAACAAAT CCTAAAATTC	840
	AGAGCATCTA TTTGGAAAGG TTTCCAATCT TTAAGGCTTG TTTTAGTGCT AATGGGGAAG	900
	AAGTTTTAGC CACGAGTACC CACAGCAAGG TTCTTTATGT CTATGACATG CTGGCTGGAA	960
50	AGTTAATTCC TGTGCATCAA GTGAGAGGTT TGAAAGAGAA GATAGTGAGG AGCTTTGAAG	1020
	TCTCCCCAGA TGGGTCCTTC TTGCTCATAA ATGGCATTGC TGGATATTTG CATTTGCTAG	108
55	CAATGAAGAC CAAAGAACTG ATTGGAAGCA TGAAAATTAA TGGAAGGGTT GCAGCATCCA	114
	CATTCTCTTC AGATAGTAAG AAAGTATACG CCTCTTCGGG GGATGGAGAA GTTTATGTTT	120

GGGATGTGAA CTCAAGGAAG TGCCTTAACA GATTTGTTGA TGAAGGCAGT TTATATGGAT

PCT/US98/11422

	TAAGCATTGC CACATCTAGG AATGGACAGT ATGTTGCTTG TGGTTCTAAT TGTGGAGTCG	1320
	TAAATATATA CAATCAAGAT TCTTGTCTCC AAGAAACAAA CCCAAAGCCA ATAAAAGCTA	1380
5	TAATGAACTT GGTTACAGGT GTTACTTCTC TGACCTTCAA TCCTACTACA GAAATCTTGG	1440
	CAATTGCTTC AGAAAAAATG AAAGAAGCAG TCAGATTGGT TCATCTTCCT TCCTGTACAG	1500
10	TATTITCAAA CTTCCCAGTC ATTAAAAATA AGAATATITC TCATGTTCAT ACCATGGATT	1560
10	TITCTCCGAG AAGTGGATAC TTTGCCTTGG GGAATGAAAA GGGCAAGGCC CTGATGTATA	1620
	GGTTGCACCA TTACTCAGAC TTCTAAAGAG ACTATTTGAA GTCCAGTTGA GTCACAAGAG	1680
15	AAGCCTGTCT TGATATATCA TCTCAGAAAC TTTCCTGAAT ATGTGATAAT ATATGGAAAA	1740
	TGATTTATAG ATCCAGCTGT GCTTAAGAGC CAGTAATGTC TTAATAAACA TGTGGCAGCT	1800
20	ТТТСТТТСАА АААААААА ААААААААА ААААААААА АААСТССА	1848
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# (2) INFORMATION FOR SEQ ID NO: 189:

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# (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1146 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- 30 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 189:

AAAAAAAACC CAGGGGAACN TTGGGGGCCG CTTTNNNTTC CCCCTCCAGG CCATTGGGGA	60
ATTCTTCAAG TTAATCCTGC TTTGCTCTTG GCCAACAGGG CTTGTAGGGG GGAGAGACCC	120
AGGATCATCA AGGGGTTCGA GTGCAAGCCT CACTCCCAGC CCTGGCAGGC AGCCCTGTTC	180
GAGAAGACGC GGCTACTCTG TGGGGCGACG CTCATCGCCC CCAGATGGCT CCTGACAGCA	240
GCCCACTGCC TCAAGCCCCG CTACATAGTT CACCTGGGGC AGCACAACCT CCAGAAGGAG	300
GAGGGCTGTG AGCAGACCCG GACAGCCACT GAGTCCTTCC CCCACCCCGG CTTCAACAAC	360
AGCCTCCCCA ACAAAGACCA CCGCAATGAC ATCATGCTGG TGAAGATGGC ATCGCCAGTC	420
TCCATCACCT GGGCTGTGCG ACCCCTCACC CTCTCCTCAC GCTGTGTCAC TGCTGGCACC	480
AGCTGYCTCA TTTCCGGCTG GGGCAGMACG TCCAGCCCCC AGTTACGCCT GCCTCACACC	540
TTGSGATGCG CCAACATCAC CATCATTGAG CACCAGAAGT GTGAGAACGC CTACCCCGGC	600
AACATCACAG ACACCATGGT GTGTGCCAGC GTGCAGGAAG GGGGCAAGGA CTCCTGCCAG	660
GGTGACTCCG GGGGCCCTCT GGTCTGTAAC CAGTCTCTTC AAGGCATTAT CTCCTGGGGC	720
CAGGATCCGT GTGCGATCAC CCGAAAGCCT GGTGTCTACA CGAAAGTCTG CAAATATGTG	780
GACTGGATCC AGGAGACGAT GAAGAACAAT TAGACTGGAC CCACCCACCA CAGCCCATCA	840

	CCCTCCATTT	CCACTTGGTG	TTTGGTTCCT	GTTCACTCTG	TTAATAAGAA	ACCCTAAGCC	900
5	AAGACCCTCT	ACGAACATTC	TTTGGGCCTC	CTGGACTACA	GGAGATGCTG	TCACTTAATA	960
	ATCAACCTGG	GGTTCGAAAT	CAGTGAGACC	TGGATTCAAA	TTCTGCCTTG	AAATATTGTG	1020
	ACTCTGGGAA	TGACAACACC	TGGTTTGTTC	TCTGTTGTAT	CCCCAGCCCC	AAAGACAGCT	1080
10	CCTGGCCATA	TATCAAGGTT	тсаатааата	TTTGCTAAAT	GAAAAARAAA	AAAAAAAAA	1140
	ACTCGA						1146

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#### (2) INFORMATION FOR SEQ ID NO: 190:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 906 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 190:

ACTCCCTCAC CCAGGTCCCA GCCCTGGGAA CCACCTACCG TGAGCCCTTT TGCAGATATA GACTCATTTC ATCCTCAGAT GGTCCTTCAA GGTAGGTACT TTAGTCCCAT TTTAGAGATG 120 AGACGATTGA GGCCAGAGGG GTGNNGTAAC TTGCCTGGGG GCTCACGAGC ACAAAAGGAG 180 CCGAGGCAGG ATCTGACCCT TGTTCTCTGG CCTCACTGCC CTCACTTTGC CATGACCCGA 240 AGITATGTCC CTACAAAGCA ATGCATGGTC CAAGGYTCTT TTTATTGTAT TTTTATTTTT 300 AAGGGTCCTG TTCAAAACTG GTGTGAGCTC TGAGGAGTCC TGAACCCTGG GTGCAGCATC 360 CTAGCATCCT GGGAGTCCTT TTCTGCCCAC ACTGAGCTGG GCTCCTCGAG GGGTGGGGCT 420 GCTGTCCCTG GAAGCCTGGC AGCAGCACTG TATCGGGTTG GCTGAAGCTG ARCGCCGTGG 480 GGTGCAGGGC TCCMGGAATC CCCGTTTGGC TGAAGGGGTT CCCTGTAGCC MGGGATGTTT 540 ATGAGGTCTC TCTGATGCCC CAGGCGCAGG ACATGTGTGC GGGTGGAGAA AAGCAGGCCC 600 TITCAGTGCC AGCTCCACTC AATTTCTATG TGGACCAAGA ACGATAAACT TAAAAAATTT 660 TTTTTCCTAA GGTATCTTCA GAATATGGTG TATTTTTATG TGGAAAAGAA AAGTTATGAA GGCAGCTGTT ACTITAAGAG AAAATTCATT AAAAGTCCTC GAGGTATGAA GATGACGGCG 840 CAAGCATGTC AGACAATAAA GTCTTTGTAA AAAGRGAAAA AAAAAAAAA AAAAAAAAA 900 906 ACTCGA

#### (2) INFORMATION FOR SEQ ID NO: 191:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1941 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 191: 10 CTTCAGCTGA AGCCCAGGGA CCCCTTTTCC ACCCTGGGCC CCAATGCCGT CCTTTCCCCG 60 CAGAGACTGG TCTTGGAAAC CCTCAGCAAA CTCAGCATCC AGGACAACAA TGTGGACCTG 120 15 ATTCTGGCCA CACCCCCTT CAGCCGCCTG GAGAAGTTGT ATAGCACTAT GGTGCGCTTC 180 CTCAGTGACC GAAAGAACCC GGTGTGCCGG AGATGGCTGT GGTACTGCTG GCCAACCTGG 240 CTCAGGGGGA CAGCCTGGCA GCTCGTGCCA TTGCAGTGCA GAAGGGCAGT ATCGGCAACC 300 20 TCCTGGGCTT CCTAGAGGAC AGCCTTGCCG CCACACAGTT CCAGCAGAGC CAGGCCAGCC 360 TCCTCCACAT GCAGAACCCA CCCTTTGAGC CAAYTAGTGT GGACATGATG CGGCGGGCTG 25 CCCGCGCGCT GCTTGCCTTG GCCAAGGTGG ACGAGAACCA CTCAGAGTTT ACTCTGTACG 480 AATCACGGCT GTTGGACATC TCGGTATCAC CGTTGATGAA CTCAKTGGTT TCACAAGTCA 540 TTTGTGATGT ACTGTTTTTG NATTGGCCAG TCATGACAGC CGTGGGACAC CTCCCCCCC 600 30 CGTGTGTGTG TGCGTGTGTG GAGAACTTAG AAACTGACTG TTGCCCTTTA TTTATGCAAA 660 ACCACCTCAG AATCCAGTTT ACCCTGTGCT GTCCAGCTTC TCCCTTGGGA AAAAGTCTCT 35 CCTGTTTCTC TCTCCTCCTT CCACCTCCCC TCCCTCCATC ACCTCACGCC TTTCTGTTCC 780 TTGTCCTCAC CTTACTCCCC TCAGGACCCT ACCCCACCCT CTTTGAAAAG ACAAAGCTCT 840 GCCTACATAG AAGACTTTTT TTATTTTAAC CAAAGTTACT GTTGTTTACA GTGAGTTTGG 900 40 GGAAAAAAA TAAAATAAAA ATGGCTTTCC CAGTCCTTGC ATCAACGGGA TGCCACATTT 960 CATAACTGTT TTTAATGGTA AAAAAAAAA AAAAAAATAC AAAAAAAAT TCTGAAGGAC 1020 45 AAAAAAGGTG ACTGCTGAAC TGTGTGTGGT TTATTGTTGT ACATTCACAA TCTTGCAGGA 1080 1140 GCCAAGAAGT TCGCAGTTGT GAACAGACCC TGTTCACTGG AGAGGCCTGT GCAGTAGAGT GTAGACCCTT TCATGTACTG TACTGTACAC CTGATACTGT AAACATACTG TAATAATAAT 1200 50 GTCTCACATG GAAACAGAAA ACGCTGGGTC AGCAGCAAGC TGTAGTTTTT AAAAATGTTT 1260 TTAGTTAAAC GTTGAGGAGA AAAAAAAAA AGGCTTTTCC CCCAAAGTAT CATGTGTGAA 1320 55 CCTACAACAC CCTGACCTCT TTCTCTCCTC CTTGATTGTA TGAATAACCC TGAGATCACC 1380 TCTTAGAACT GGTTTTAACC TTTAGCTGCA GCGNCTACGT CNAWCGNIGT GTATATATAT 1440 GACGTKGTAC ATTGCACATA CCCTTGGATC CCCACAGTTK GGTCCTCCTC CCAGCTACCC 1500 60

	CTTTATAGTA	TGACGAGTTA	ACAAGTTGGT	GACCTGCACA	AAGCGAGACA	CAGCTATTTA	1560
ہے	ATCTCTTGCC	CAGATATCGC	CCCTCTTGGT	GCGATGCTGT	ACAGGTCTCT	GTAAAAAGTC	1620
5	CTTGCTGTCT	CAGCAGCCAA	TCAACTTATA	GTTTATTTTT	TTCTGGGTTT	TTGTTTTGTT	1680
	TIGITITCIT	TCTAATCGAG	GTGTGAAAAA	GTTCTAGGTT	CAGTTGAAGT	TCTGATGAAG	1740
10	AAACACAATT	GAGATTTTTT	CAGTGATAAA	ATCTGCATAT	TTGTATTTCA	ACAATGTAGC	1800
	TAAAACTTGA	TGTAAATTCC	TCCTTTTTT	CCTTTTTGG	CTTAATGAAT	ATCATTTATT	1860
٠, -	CAGTATGAAA	TCTTTATACT	ATATGTTCCA	CCTCTTAAGA	ATAAATGTAC	ATTAAATCTT	1920
15	GGTAAGACTT	тааааааааа	A	•			1941

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#### (2) INFORMATION FOR SEQ ID NO: 192:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2118 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 192:

AAATAATAAT AANAATAAAT AAAAATWAAG TGCTTAKTGT AACTCAGCGG ACAGGGCTCC 60 CAGCTGCTCT GGCACGTGGG ACACCYTCCA CCCTGCACAC AACAGGCATG CAAAGAGGAC 120 TOGATATGCT GGGGTAGAGT GCTTCTGGTG TGTTCACTTT AAGAAAACAT CTGCCAAGAG 180 35 AGAAGAGTGC CCAGGAAAGA CCAGGAAAAT ACAAGTACAT GGCTGCTTCA TACCATATAC 240 CCCAATTCTT TAAAGCAGCA AAAGGCACTT TTTTTTCAG GCCAGAGTGA ATCTAAAACA 300 40 AACCTGGCTT TGCTTACAGG GAAGCTGTCC CAGAAGGACT GAGTGATGCC TCTTGTTCCC 360 TAAGGTCTGG AGAGTCTTTG CAAGTTTCCA ACGACATTTC CAACCAGGTG GGAGAGACCA 420 GCAGTTGACG AGACAAGTCA GACCCAAAAA ACGACGCCAA GGTAGTGAGT GGGTGCCTAT 480 45 TTGGGAGTAG GATGATTTGA GGAAAACAGG AAGAAAAACC GGTCAGAAAG TGGCACTTTG 540 GAAGTGGAAA GCTGTTTGCA AATAGCAACT CTGGCTAAAG CGAAAATGTT AATCAAGTAG 600 50 AAAGTAAAAT TCAGGATCTT AGAAGCTCAT CCTTCTGATG AGAACTATTT TTTTTTCCGT 660 GAAGGAACTA TTATTACTTT AAAAGTGAGG GTAATTTACA TATGGGGTGT ATATATTCTA 720 AAAATAGTAA TAAAAGTACC TTTTATAAGC AATGTTGTGT GGCTTGTAGA AGAAAGCAGG 780 55 GAGGAAAAAA AGGCAGGCAA AACTAGTCTA GGTCTAGGCC CTAAAAAATGA GCTTCCTTCC CACTIGACIG GAAACGCCCA TGIGATITCT AGGCTGAAAA TAGGTAGGAT TTAACGAGTA 900

	ACCTAGTTCC CTTCTGTCTC TGATTTCTGA TCAGCTGATG GAGCTGCTAG TAAGAGGGGC	960
	CGATCATGCT CCCAGACGAG TCCTTTGGCC TCTTGCTCTC CATCCCAAGC CTGACTCCTT	1020
5	CAGCAGCAGC CCCCTCCTTC TGTGTCCATC TGATGCAGGC AAGCAGGAGC AGTAAGAGGG	1080
	CATCCCATGT TCCAGTTCAC CTTCTATGGG GTGACTARGA GGTTCCCGGT AACTAGGGCA	1140
	GCCCARGCCC AGCAGGTTGC AAAAGCAGCT GCAAGCTTCA GAAACCCACT TCCTCCAACA	1200
10	CCAGGGAGGT GGCAGAGAGC CCATCCAAAA GCCCACTGGG AGAGGCATAA GATTCTGTGC	1260
	CAGGCCCCCA GGTCCCCTCT GTGTCAGGTA GGCTCTGCTA CTGGCCTCTG AAGTAAAGGC	1320
15	AAANACAAAC GGGCAGGGCA GGGTGGCAGG AATAAAAAAC TCTGGACAGA AACCCTTTTA	1380
	ATAAAGGAAA TTCCACCCCT CCCAATCCTT CCATGGAAGG GTGAGACCTT AATGTGATGT	1440
	AAGAGGAAGG TCTTCTCTGG CTTTCAGGGA AACAGCTGCA GCTGAAACTT AGGGGCCCAT	1500
20	TCCAGGGCAC TTTTCACCAC AGCCAGTGCA GCCGCTCCAA GTGCCACTGT CAGCCCCATC	1560
	ACTGCCAATT TCACAAAGCG GTTGGTCCTT GGCTTGGTCA GGACATCTTT TGTTCGATCT	1620
25	TCAGGCCGCA GAAGTCCCCG AANACCGCTG CCGCAGCACC ATATCAGGCC TCTGCTGGGC	1680
	TGATGCCAGC TCAAAGTCTT TGAAAGTAGA GGCTGCCGTC CTCTCAGCTT GCTGTTGGGC	1740
	AGCGGCCTCC CGAGCAAGTT CGGATGGGGG AAACTGAACA AAAAGGTCTC CTSTCTGCTG	1800
30	ATCAGTGTCT CATAGGGCAA GTCCTGAGGG ATCTGGGACA ACAGGTGGTG GACCGAGGCC	1860
	ATGTCACAGT CACAGTCCAG GACTTCCTGC TCGCGATACA ACACAATCAC GGCTGCAAAG	1920
35	TAAATCGGCA TCAGTGGGTG GCAGGCCAGG AAGAAGTCAT ATAACCGCAC GACGTGCCTG	1980
	AAGTCAGACA GGACATGCCC AAACCAGGTG ATGAGCCAGC TGAGGGCAAA GATGGTCCCT	2040
40	ACCTCAGCAC TCTGCATGAA GTCATGGAGC TCTGGATTCA CCTGGTCAAT GATGGGCATC	2100
40	AGATAGTTTA ATATATCC	2118

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# (2) INFORMATION FOR SEQ ID NO: 193:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1538 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 193:

CCGGGTTCGG CTCTGTGTCA GCAGCCGGGC GGCGCTCGGG CGGGACATGG CAGCCTGTAC 60

AGCCCGGCGG CCTGGCCGTG GGCAGCCGCT GGTGGTCCCG GTCGCTGACT GNGGCCCGGT 120

60 GGCCAAGGCC GCTCTGTGCG CGGCCGNAGC TGGAGCCTTC TCGCCAGCGT CGACCACGAC 180

	GACGCGGAGG CACCTCTCGT CCCGAAACCG ACCAGAGGGC AAAGTGTTGG AGACAGTTGG	240
_	TGTGTTTGAG GTGCCAAAAC AGAATGGAAA ATATGAGACC GGGCAGCTTT TCCTTCATAG	300
5	CATTITIGGC TACCGAGGTG TCGTCCTGTT TCCCTGGCAG GCCAGACTGT RTGACCGGGA	360
	TGTGGCTTCT GCAGCTCCAG AAAAAGCAGA GAACCCTGCT GGCCATGGCT CCAAGGAGGT	<b>420</b>
10	GAAAGGCAAA ACTCACACTT ACTATCAGGT GCTGATTGAT GCTCGTGACT GCCCACATAT	480
	ATCTCAGAGA TCTCAGACAG AAGCTGTGAC CTTCTTGGCT AACCATGATG ACAGTCGGGC	540
15	CCTCTATGCC ATCCCAGGCT TGGACTATGT CAGCCATGAA GACATCCTCC CCTACACCTC	600
15	CACTGATCAG GTTCCCATCC AACATGAACT CTTTGAAAGA TTTCTTCTGT ATGACCAGAC	660
	AAAAGCACCT CCTTTTGTGG CTCGGGAGAC GCTAAGGGCC TGGCAAGAGA AGAATCACCC	720
20	CTGGCTGGAG CTCTCCGATG TTCATCGGGA AACAACTGAG AACATACGTG TCACTGTCAT	780
	CCCCTTCTAC ATGGGCATGA GGGAAGCCCA GAATTCCCAC GTGTACTGGT GGCGCTACTG	840
25	TATCCGTTTG GAGAACCTTG ACAGTGATGT GGTACAGCTC CGGGAGCGGC ACTGGAGGAT	900
	ATTCAGTCTC TCTGGCACCT TGGAGACAGT GCGAGGCCGA GGGGTAGTGG GCAGGGAACC	960
	AGTGTTATCC AAGGAGCAGC CTGCGTTCCA GTATAGCAGC CACGTCTCGC TGCAGGCTTC	1020
30	CAGTGGGCAC ATGTGGGGCA CGTTCCGCTT TGAAAGACCT GATGGCTCCC ACTTTGATGT	1080
	TCGGATTCCT CCCTTCTCCC TGGAAAGCAA TAAAGATGAG AAGACACCAC CCTCAGGCCT	1140
35 -	TCACTGGTAG GCCAGCTGAG GCCCCAAGTG CCCAGGCTTG GTCACCGGGA AGAACAACTC	1200
-	TCATCCCACA ATTGCTGCAG AACTCTTCTC TCCCCATCAT GGGCCACAGT GGGTCTCTTA	1260
	ATTIGATIGI GGGGTTCTTT TTGTGGGGAG GGGTGGTATA ACTITTCTTC AGAAGACCCA	1320
40	TGTGGGACAC CTCCAAGGCT GGCCTCCTCA TAAGCCCTGC CTACACCATG TTCCAGTAAA	1380
	CCTCTCCACC AAGGAACTGT GTTCAGCTGC CACAGGCCTG GAGGAGTTTC CTGGCCTGTC	1440
45	ACGTGAGGTT TGATCAGTAA ACCAGTGCAS GYTTGGCCAA AAAAAAAAAA AAAAAAAAA	1500
73	АААААААА АААААААА ААААААААА АААСТССА	1538

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(2) INFORMATION FOR SEQ ID NO: 194:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1098 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 194:

-	AGACCCTGTC TCAAATAATA ATAATAATAA TAATCTTATT TTGGAGAATA AAGAGACC	rs 60
	TGGATTTGAG GTGCCATTTG GGTAGAAAGA AAAGACGTTT ACACCGAGAA ATAGTCTG	rg 120
5	TTGCCCTGAA GGAGCAGAGG GATGCATCGC TGGAGGTGAC CTACAGTTGA AGAAGACT	CA 180
	TTATGACAGA CCTTGTCCTT CTTCCTTGTG GAAAGTGTTT CCTCTGCTGC TACTGCTC.	AT 240
10	GAGACTCTTC CCCCTCCCTG TCCCAGGGAA CCAAAGGGCT TTNCTACCAC ACCCTTTC	rr · 300
10	NGCCCCCCGC CTCCCATGTC TGCTGTGCCT TTGTACTCAG CAATTCTTNG TTTGCTCC	CA 360
	TTATCTTCCA GCCGGATACA GAGTGAATAG TTAACCACAC TTAGGTCAAA TAGGATCT	AA 420
15	ATTITITGITC CIGCICCNGT GIAAAGAGGC CAGIGITITGI GIGITGCAAG CAGCCTIG	GA 480
	ATAGTAACTC TTCTCATTTG TTTGGGATCT GGCCAMCAAG TTCCAGAATG ATACACGG	AT 540
20	CAGTGCAGAA GTTCATCAGG CTCTCGGACC TTAGGGCTGT TGGAGAAGGC TTCAGCAG	CA 600
20	GAACTGATGG TKAWKGYTCG TGTTCTCCAT CCTCAACTTT CTTTGCTTCG ATCATACA	.CA 660
	AGAATACATT TGGAAGGGCA AAAAATGAAC ACTGTTGTTC ATTGCAGCCG TGTTTTGT	GA 720
25	CACAGATGCA CAGTCTGCTG TGAAGACCTT CTCTCAAGTG GSATYTGGGA GTCCATGC	CA 780
	GATCATGGTG CTTCATGAGA GACTGACAGC TATCAGGGGT TGTGGCACTT AGTGAGGA	ACT 84
30	CTCCTCCCCC AGTGTGTGCT GATGACACAT ACACACCTGA CAATAGCTTG AGTCTTCT	rcr 90
50	GTTCCTTTTA CTCTGTAGCC AACATACACA TGATTTAAAA CCCTTTCTAA ATATCTA	rca 96
	TGGTTCATCC TTGTCCAAAT GCAGAGTCAG AGCTATTTGT ACTTCATTAT TATTTCCA	AAG 102
35	GCGAATAGIT GGCTTTCTTT TIGCAAAAAT AATTAAAGTT TITGTATGTT GCAAAAA	AAA 108
	AAAAAAAAA CTACGTAG	109

#### (2) INFORMATION FOR SEQ ID NO: 195:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1001 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 195:

	GAATTCGGCA	CGAGATAGCT	TGCATCTCAT	CCCAGTAAAA	CCACTTATTT	ATAACATATC	60
55	AACGTATTGA	CAAGGTTGAA	GAGCAAGATT	GTTCTGAGGT	GAGATGCAAA	TTTCAAAGGG	120
	GTGAGCACTA	ATTGTTCCAG	TGATTGTTTA	TTTATTGGCT	AGGACATAAT	TACTCTCTTT	180
	GAGGTTACAC	ATCTGCCTCC	AGGTTCCTGT	GTGCTTGTGC	CCTTGGGATC	AGGCCAGGGC	240
60	AGACTGTGAT	CACTGAGATT	CAAACTCCCA	GARTAATCAG	CAAGAGCTTT	CTAGAGACCA	300

	AGGCCAGGCC TGATCCCTGA GGGATGCATG AGAAGGCTTG GAATCTCATT CTGCTATGGT	360
_	GGCTCTCTCT TGATCTTCTT GGAGTAGCAA AAACAGCAAT GTGGGCCCCAA TGGTGTGGCC	420
5	TAAATGATCA CAAAGGTAAA TGAGTAAAGG GCTCAGCAGA TGAGTAAGGA GCCTTGTCCT	480
	GAGAAATTAG CACTGGGCTC TGCATTCAGA AACATGTGAT AAGCATTGCC CATTGCACAT	540
10	TGCCTTTATT GTGTAAGGAC ATGAAATTCC AGTTTTGCAT AGCTAGTGAT GAATACCTGA	600
	AGGGAATTGC AGACATATTT TATTTTATTT TTAATTGACA GATGGAATTG TATATATTTA	660
. ~	TCATGTACAT AATCATGCTT TAAAATATGT ACATTATGGA ATGGCTAAAT CAAACTAACC	720
15	TAGGCATTAT CTCATATAAT TGTCATTTTT GTGGCGAGAA GACTAAAAAT CTACCCTTTC	780
	AGCATTITTA AAGAATACAA TGTGTTTTAT TAACAACAGT CACCATTTGG TACACTAGAT	840
20	CTCTTGAACT TCTTCCTCTT ATCTAACTGA GATCTTGTAA CCTTTGATAA CAGCTCCCAA	900
	GCCCTTCCCC AACCACTGCT CCACCCGTGG TAACCACCAT TCTATTCTCA ACTTCCTGGT	960
05	AATCACCATT CTAGACACAG GGAAGACTCT CTACCCTCTG A	1001
25		
	(2) INFORMATION FOR SEQ ID NO: 196:	•
30	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 1443 base pairs (B) TYPE: nucleic acid	
35	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
33	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 196:	
	ATAAACTGAA ATAGGTCATG CAAATATAAA ATATTATTT TAAATTATTT GTCATAAGAA	. 60
40	ACGATGGTGG CCATATTTTG CTTTAATAAT GGAAAAAATG TGGTTAGCAT TCTKTGGAAG	120
	GTGGTCATCA GATAGTAGAC ATTTTCTAGG ATTTATTTCT ACCTGCATAT GTGGAAATGT	180
45	GTACTACTTT AGATTTATWT AATGGCAGCT AACTCAGAGG CATCAAAATG TGCTAATGGT	240
,73	GTACTACTT AGAITTATUT ANGOCASCT TOTAL TOTAL COLORS ARGGECAGGG	300
	CCGTACAGTG AACTTGTCCT TTGSCAGACG CCAGCGTCTG CCCCTGACCC CGTCTCCACT	
50	CTCTGTGTCC TGGAGGAGGA GCCCCTTGAT GCYTACCCTG ATTCACCTTC TGCGTGCCTT	
		480
55	GTACTGAACT GGGAAGAGCC GTGCAATAAC GGATCTGAAA TCCTTGCTTA CACCATTGAT	
55	CTAGGAGACA CTAGCATTAC CGTGGGCAAC ACCACCATGC ATGTTATGAA AGATCTCCTT	

TGGGGATCTA AGTAAACCTC TCGGGGAAAA TGACCAAGTG GATGTCATCT CCCAGCTGTT

	TCTAAGAGCC	CAGATGTCCA	GAGTATTGTC	TCACCTTGAT	CCCTCAGGCC	AGAAGACCTG	720
	TGAAAAAGCC	ACACTGGTTC	AGGGACTCAC	TGGACGGTTT	TGTGTCCACT	YTAXOTTGCA	780
5	CCGTCTCTAC	CCCAGAGTGG	ACTCARATCC	TCAAGTCATC	CTCTGAACAT	TGREGTCAGA	840
	AATTATAAAA	GGGCTTTGGC	AATATGTTAG	CCCAAGAATT	TECCTTOTTE	CAGAAATTGT-	900
0	GCCGACNITA	ACAGTGGCTT	AAATGATGGT	AAAACTTTTA	AGATTTCTAA	AAGGRTGGCA	- 960
U	TTGGAGATAC	GTTGACTTTT	ATTAAACMAC	CTATAGTTGT	TTAATGAYTT	CTAAAAAAT	1020
	ATCTGGAGCT	CAGGGGTTCA	ACTGAGGGAA	CACATGITGA	GRATCATTGT	TTACTAATTA	1080
15	AATGCCAGGT	AACCCGTTGA	AATTATCAAA	AACATOTTOC	ACGTACCAGA	AAGGACOTCA	1140
	GAGGATAGTT	CTGTTATGGA	GAAGATGAAA	. TGGTTTAGTA	GTGTAGGAAC	TATGGAAAGG	1200
20	TGAGCTTAGA	TTTGGATAGT	AAAACCTCAA	GACCOTACTT	AAAAAGTATT	TTATGAATGC	1260
20	AGCATAAATA	ATTTAATTCA	GTGTTAANAT	GCCAAGGCTA	GTATATTGAG	CTGAATGTGA	1320
	AAAGAAACTC	ACATTGGGAG	AATGCCACCT	TITCCTIATA	AGATAGCTTT	GAASATACCA	1380
25	TTTTAGACAG	ATGGAAATTG	AATAGCTTTA	GAAAAGGCAA	ATGTTTGATO	TTGGGGAAAA	1440
	AAA						144

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# (2) INFORMATION FOR SEQ ID NO: 197:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1282 base pairs

- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: dcuble
- (D) TOPOLOGY: linear

40 (xi) SEQUENCE DESCRIPTION: SEQ D NO: 197:

		=					
	GAAAAAAAA A	AGTATGACCC	AGTAGCTAGG	CYCCLELECC	CCCGCCAAGT	TGACACATAA	60
45	AATTAACTGT (	CACAGTATCA	TCTTAGAAGT	CAAACAAGCC	CCTTTATCCT	GCAGTGCCCC	120
43	TCTACCACCA (	CCTACTGACA	AAGAACATGG	TGCTATCTGG	CĄTGGGAGAA	AUGUTCAGTT	180
	TGCTATGGCT T	IGTATGTGTC	CCCTCAAATT	CAAGTGTTGC	CAATGTGACA	GCATCAAGAG	240
50	GTGGGGTCTT	TAAGAGATCA	CTAGGCCATG	AGGGATTCTC	TTAGGACTGG	GATGAAGGCC	300
	САТААТАААА (	GAGGTTTCAG	GGAGCATCCT	GCTAGCTTGC	CTTCTGTATG	TGAGAACACA	360
<b></b>	GCAAGAAAGC (	CCTAGTCAAC	AAGTGCCAGC	TCCTTGATCT	TAGACTICCC	ACCOTCCAGA	420
55	ACTGTGAGAA	ATACATTTCT	GTTCCTTACA	AATTACCCAG	TCTCCTGTAT	TCTGTTATAG	480
	CAGCACAAAA	TGAAGATACC	ATACCTGAAC	ACCTGAACAT	TCTTCACAAG	GTAGTAAATG	540
60	CACTGCTTTA	TTCTGGTCTC	AGTATTGTGT	GCTTAATAAG	GAAATGAGAA	AGGSTGGATC	600

	AGGGCATAGG	ATGAACAAGT	TACTGCTAGA	CCTCTCACAA	TGCCACTAAT	GGATAAGATT	660
5	GTATTTTCAT	CATTNCTTGT	CTCTTCGGAA	GCTAACACCA	TGCTATAATA	GGCADTAAAT	720
3	AGATGTCTAA	AAACACCTTA	AGTATTIGIC	TAGAAATOTG	GTGCATTGTC	CAGAAAGAAC	780
	CAAAATTCMA	AATAATTTCA	AAGGCCTAA	AGCACTAXIT	ANTOMANT	CATTAGTTTT	840
10	TAATGGTACT	ACCACTCTCA	TAAAATTTAA	GTCATCTTAC	GIICCICIIC	CTCGCATTGG	900
	ATTTATTGCT	AAAACCTGGT	AAACACTTTA	ATCCYTTICA	ATTCCATTAC	CACTGCTCTT	960
15	GTCCAGAATT	ACTCGCAGAC	TAATAGTCAC	CIGACTICIS	CCCCTGCATC	CCGAITTGCT	1020
13	GTCTAATTCT	GGTTACAAAT	AAGTAACTGC	CAAACTAATC	TTTCTAAAAA	GCAAGACTGA	1080
	TCTCGTCACT	CCTTTGCTCA	ACAATGTAAA	AGCTCCCATT	GTCTCCCAAA	TARRACCAGO	1140
20	TTTCCACTGT	GTATACAATA	CATCCATGAT	CTGTATCCAG	CATCATTTTG	TATTIGCTCA	1200
	CTTTATACAC	CACCCCCAT	GCCACATCAA	. ATTAAATTAT	CCTGATAAAT	GCAACTGCAA	1260
25	<b>АААААААА</b>	AAAAAAACTC	GA				1282

# (2) INFORMATION FOR SEQ ID NO: 198:

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## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 951 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 198:

40	ATTTCGGAAC	GAGGACTGAA	GTGGGAGCGG	CGGCAGGGTA	GAAGACAGAA	GGGGGATCTA	60
40	TGTGGTAACT	AAAGAATGTT	TCTCTTTTGT	TAATTATTGT	GEGIGEGE	TTTTATTGTT	120
	TGCTTAAGAG	AATCAAAAAC	TGAAAAAAAT	GAGAATACAG	GAAATGGCTC	TIGITITATIT	180
45	TTTTGCTGTG	TTTACAGCTT	GTTAATGCTC	TACTGTCTTT	GTTTCAAGAG	AGATTTGTTC	240
	ACTGCCCAGC	TCGTTTTGTG	TCCTGAGCCC	TATGCCCAGC	CCACCTTATA	AATCATGCCT	300
.50	GTTTAGATGT	TTGATTTTGT	TCTGTTTGCT	ATTGTTATCT	TAAAGGTGTA	TAACTCTGAC	360
50	ATGCCAGACA	TCAAATTAAG	СТСАААТТАА	GCTCTCGTTT	ALATGITTAA	ACACCTAATT	420
	TATATTCTAA	TTGATCCCAG	CCACTGATGC	ATGTACTTTA	GCTACTTCTG	CTARATAAGC	480
55	ATATTAATTT	TCCACATCAG	GCCATCAGAT	CTTGAGAACC	AACAGTTATC	TAGAATTCCG	540
	TGTCTACTAA	TGTTTCACCT	GCATGCAGCC	TTCATTAATT	TTGTAGCAAA	ATATAAAGTG	600
60	ATCATTATGT	AGTITCIGGA	TAAAAAATT	TTGTGTGTGA	AGTTGCTTTG	TAAAGTGCAT	660

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	GTGGAATTAA TGGGACAGTG TGCCCTTTGT GTTAGATGTT AGAGCAAAAG AAAGGGCTTA	720
	TAGTGTTAGT ATTGGAGCAC TTTGAAGATA GATATTTTCA GAAAAGATGT AGGATTTAAA	780
5	AGTTAAATTT TAAATTTTAG AAAAAGATAT GATGGCAATT GGAAATAGTC ACAATGAAGT	840
	TCTTCATCCA GTAGGTGTTT AACAGTGTTA TTTTGCCACT GGTAATGTGT AAACTGTGAG	900
	TGATTTACAA TAAATGATTA TGAATTCAAA AAAAAAAAAA	951
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15	(2) INFORMATION FOR SEQ ID NO: 199:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1740 base pairs	
	(B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double	
20	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 199:	
25	TTATTATAAT AATGATGATG ATTCCAAGGA AAAAACCTAC AGCGAATGTT CCATTTCTAC	60
20	CCCGCACGCA GACACTCTCC CTAACACTGA TAACCTGAGC CCCCAGCACT GGACGGAAGA	120
	ATGCTGGCGT CTCCGTGTGT ACTGGTTCAG GGTTCTGGCC CCAGCCTTGT CAGGACCCCC	180
30	TGGTGTCCAG AGCCCCCACC CCTCCCGCAA CAAGCAGCTG ATGCCCCAGT GATTCTCTAT	240
	ACATTTTTCA CCTCGGCCAA TATGTCCAGG AAAACTGCTT ACTTCTCTTT TCTTGCCTGG	. 300
35	AGCCTTCATT GTTCACCCTT ACGTTGCAAT ATAGGAATTA ATGCTACAAA ATAAAAGTAA	360
33	AGCTTACCTG AAAAGTGCAT AGTTTGGGGC AATGGTATCT ACATCTCCCA CTGTGGGAAA	420
	ACCAGCAAAG CATCAAAACT CTCAATTCTC CTGTTACCRA ATGCAGATCT GAATTATAAG	480
40	ATGTTTATGT TTGACCATTG TTTCAACAAT GGGATTTTGT TACGAATTAT CCCTTTAACT	540
	GAAACCCTCA GTTTTACTGT TTACATTATT AGGAAAACAG GGATATCTTT TGAATCTAAA	600
45	AATTTGATGT ACAGCATGTG ATTTTTGAAG TTTACATGTA AAGTCACAGT ATAGGTGAAA	660
45	TAACGTTTGT CATATTTTGA GACGTATCCT GCAGCCATGT TTTTACGTGA GTGTTTTAGT	720
	CAAAGTACAT GGTAGACAGT CTTTCACAAT AAAAGGAAAA GGATTTTTTT TCCTCCAAAT	780
50	GTACATTTAT CAACCTAATG ATTGATTTTT TTAAAAAGAG ATTTCGCCCC AGTCTGGTTT	840
	ATGAAAGTTC ATTGCCCTAA ACTGTGCTGA TTGTTTTTAA TCAAGTTATA AATTTCCAAC	900
55	CTAGATCATG TATCTACCAA CTCTCCTGCA TTTTCCAAAA GGCATTGAGC TTAAATATTA	96
55	GTCTTGCTTA GAGTAGGTTA TCCACTTACA TGCTGCGCTA AAGCCATGCC TTTGAAACTC	102
	CTTGTTTAAA ACATGATATG ATTTTTGTGG GCAGTTTCAG AAAAGAAAAC AAACAAACAA	108

AAATCGACCC TTTAATTATT ACTTGCAACT CAACAGATCT CCCTGCCGTA CTGCCTTTTC

	CAGGAACTTT	ACTTCAGGGC	TGTCCAGATT	GCAGTTGTGC	CCCGTGTATG	TGGATCTAGT	1200
_	TCACAGAGTC	TTTGGAAGCC	AGCAGTCGTG	CCCTCCGTAT	ACTGTCCACT	CATTTTATGT	1260
5	AGATTTGGTA	TCCTCAGCAG	CCAGTGTTAA	CACCACTGTC	ACGTAGTTAN	CAGATTCATC	1320
	TTTTATGTAT	TTAAAGTAAT	CCATACTATG	ATTTGGTTTT	TCCCTGCACC	ATTAATTCTG	1380
10	GCATCAGATC	AGTTTTTGTG	TTGTGAAGTT	CTACTGTGGT	TTGACCCAAG	ACCACAACCA	1440
	TGAGACCCTG	AAGTAAAGAT	AAGGTACACA	TACATTATTT	GAGTAACTGT	TTCCTTGGGG	1500
15	GCCAATCTGT	GTATGCTTTT	AGAAGTTTAC	AGAATGCTTT	TATTTTTGTC	TATAACAAAC	1560
13	AGTCTGTCAT	TTATTTCTGT	TGATAAACCA	TTTGGACAGA	GTGAGGACGT	TIGCCCTGTT	1620
	ATCTCCTAGT	GCTAACAATA	CACTCCAGTC	ATGAGCCGGG	CTTTACAAAT	AAAGCACTTT	1680
20	TGATGACTCA	маааааааа	ААААААААМС	YCGGGGGGG	GCCGGTAACC	CATTINNCCC	1740

#### 25 (2) INFORMATION FOR SEQ ID NO: 200:

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#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1707 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

#### (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 200:

35	GCTTATAGAA GGGAGAGGAG CGAACATGGC AGCGCGTTGG CGGTTTTGGT GTGTCTCTGT	60
	GACCATGGTG GTGGCGCTGC TCATCGTTTG CGACGTTCCC TCAGCCTCTG CCCAAAGAAA	120
40	GAAGGAGATG GTGTTATCTG AAAAGGTTAG TCAGCTGATG GAATGGACTA ACAAAAGACC	180
40	TGTAATAAGA ATGAATGGAG ACAAGTTCCG TCGCCTTGTG AAAGCCCCAC CGAGAAATTA	240
	CTCCGTTATC GTCATGTTCA CTGCTCTCCA ACTGCATAGA CAGTGTGTCG TTTGCAAGCA	300
45	AGCTGATGAA GAATTCCAGA TCCTGGCAAA CTCCTGGCGA TACTCCAGTG CATTCACCAA	360 -
	CAGGATATTT TTTGCCATGG TGGATTTTGA TGAAGGCTCT GATGTATTTC AGATGCTAAA	420
50 .	CATGAATTCA GCTCCAACTT TCATCAACTT TCCTGCAAAA GGGAAACCCA AACGGGGTGA	480
30	TACATATGAG TTACAGGTGC GGGGTTTTTC AGCTGAGCAG ATTGCCCGGT GGATCGCCGA	540
	CAGAACTGAT GTCAATATTA GAGTGATTAG ACCCCCAAAT TATGCTGGTC CCCTTATGTT	600
55	GGGATTGCTT TTGGCTGTTA TTGGTGGACT TGTGTATCTT CGAAGAGTAA TATGGAATTT	660
	CTCTTTAATA AAACTGGATG GGCTTTTGCA GCTTTGTGTT TTGTGCTTGC TATGACATCT	720
60	GGTCAAATGT GGAACCATAT AAGAGGACCA CCATATGCCC ATAAGAATÇC CCACACGGGA	780

	CATGTGAATT ATATCCATGG AAGCAGTCAA GCCCAGTTTG TAGCTGAAAC ACACATTGTT	840
	CTTCTGTTTA ATGGTGGAGT TACCTTAGGA ATGGTGCTTT TATGTGAAGC TGCTACCTCT	900
5	GACATGGATA TTGGAAAGCG AAAGATAATG TGTGTGGCTG GTATTGGACT TGTTGTATTA	960
	TTCTTCAGTT GGATGCTCTC TATTTTTAGA TCTAAATATC ATGGCTACCC ATACAGCTTT	1020
	CTGATGAGTT AAAAAGGTCC CAGAGATATA TAGACACTGG AGTACTGGAA ATTGAAAAAC	1080
10	GAAAATCGTG TGTGTTTGAA AAGAAGAATG CAACTTGTAT ATTTTGTATT ACCTCTTTTT	1140
	TTCAAGTGAT TTAAATAGTT AATCATTTAA CCAAAGAAGA TGTGTAGTGC CTTAACAAGC	1200
15	AATCCTCTGT CAAAATCTGA GGTATTTGAA AATAATTATC CTCTTAACCT TCTCTTCCCA	1260
	GTGAACTTTA TGGAACATTT AATTTAGTAC AATTAAGTAT ATTATAAAAA TIGTAAAACT	1320
	ACTACTTTGT TTTAGTTAGA ACAAAGCTCA AAACTACTTT AGTTAACTTG GTCATCTGAT	1380
20	TTTATATTGC CTTATCCAAA GATGGGGAAA GTAAGTCCTG ACCAGGTGTT CCCACATATG	1440
	CCTGTTACAG ATAACTACAT TAGGAATTCA TTCTTAGCTT CTTCATCTTT GTGTGGATGT	1500
25	GTATACTITA CGCATCTITC CTTTTGAGTA GAGAAATTAT GTGTGTCATG TGGTCTTCTG	1560
	AAAATGGAAC ACCATTCTTC AGAGCACACG TCTAGCCCTC AGCAAGACAG TTGTTTCTCC	1620
	TCCTCCTTGC ATATTTCCTA CTGAAATACA GTGCTGTCTA TGATTGTTTT TGTTTTGTTG	1680
30	TTTTTTYGAG ATCACGYTAC TGGGCTC	1707

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# (2) INFORMATION FOR SEQ ID NO: 201:

# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 779 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 201:

CTGTCCCCAG TGTTTCCAGG TAATGACTTG GCACTCCAGA GAAAGTTTCA TRCTGTTGCG 60

TGTGGTGGCT CCAAGCCAAG CACCTGGCAT GCAGGTCAGC CCTTCCCAGC GGGCGTGGCG 120

TCGTCCTCTT CACAGATGCC ACGTTGCAGC CCCAAGGCCT CACCATTTTG CGTTTTTTAG 180

AAACCCATTT TCTTGGTCAT TTATAAAGCT GCTTTATAGA TATCTTTGAT CCTGGCATGC 240

CTTGGTTTCC TCTCCCTTCC CTCTTTCCAA TCCTGGTTTC CTAACCTCCT CTTGTAGTAA 300

TTCTCAACTC AACTCAAAGT CCCAAGAATT TGGAATGGTA GGATGCTGTG CGGGGAGCTC 360

GAGGCTGAGG CATAATCACT GCTTCGGTTC TGCTCATCAG GGGACACGCT CCCTTACTCA 420

TGGCAGCCAT GTTTGATTGT CACAGAGCCC CCCGAATACT CTGTCTATAG TGACACACTG 480

	TAGGTGTCAT	AAATTTTAAG	AAACCTGCTT	TTAAGTACTA	TTTATAGGTT	TTTCTGTTAT	540
5	ACTTGCAACC	TAGTTTTAAA	ATACATGAGG	ATTTTATGAA	AGCTTTATAC	AGACATTTAT	600
	AGGAAACTCA	TTCTTTGATT	TTAGGTGCCA	TTTAAATTGA	TAACACTTAC	TTTATAAAAA	660
	GATGCTTTTT	GTCTGGATAG	AGCCTTATAG	TTTAAAATAT	CTTCATATAT	TGCCATTTGA	720
10	TCAAATAAAT	TTCTTACTTA	GAAAAAAAAA	АААААААА	ААААААААА	AAAACTCGA	<b>7</b> 79

#### 15 (2) INFORMATION FOR SEQ ID NO: 202:

# (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1617 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 202:

	· · · · · · · ·	
25	GGCACAGCTT TCTGTCTCTT CCTCGCTCCC TCTCTTTCTC TCCTCCCTC	60
	TGCATAAAGT CTCTGTCGCT CCCGGAACTT GTTGGCAATG CCTATTTTTT GGCTTTCCCC	120
30	CGCGTTCTCT AAACTAACTA TITAAAGGTC TGCGGTCGCA AATGGTTTGA CTAAACGTAG	180
30	GATGGGACTT AAGTTGAACG GCAGATATAT TTCACTGATC CTCGCGGTGC AAATAGCGTA	240
	TCTGGTGCAG GCCGTGAGAG CAGCGGGCAA GTGCGATGCG GTCTTCAAGG GCTTTTCGGA	300
35	CTGTTTGCTC AAGCTGGGCG ACACATGGCC AACTACCCGC AGCCTGGGAC GACAAGACGA	360
	ACATCAAGAC CGTGTGCACA TACTGGGAGG ATTTCCACAG CTGCACGGTC ACAGCCCTTA	420
40	CGGATTGCCA GGAAGGGGCG AAAGATATGT GGGATAAACT GAGAAAAGAA TCCAAAAACC	480
40	TCAACATCCA AGGCAGCTTA TTCGAACTCT GCGGCAGCGG CAACGGGGCG GCGGGGTCCC	540
	TGCTCCCGGC GTTCCCGGTG CTCCTGGTGT CTCTCTCGGC AGCTTTAGCG ACCTGGCTTT	600
45	CCTTCTGAGC GTGGGGCCAG CTCCCCCCGC GCGCCCACCC ACACTCACTC CATGCTCCCG	660
	GAAATCGAGA GGAAGATCCA TTAGTTCTTT GGGGACGTTG TGATTCTCTG TGATGCTGAA	720
50	AACACTCATA TAGGATTGTG GGAAATCCTG ATTCTCTTTT TTATTTCGTT TGATTTCTTG	780
30	TGTTTATTT GCCAAATGTT ACCAATCAGT GAGCAAGCAA GCACAGCCAA AATCGGACCT	840
	CAGCTTTAGT CCGTCTTCAC ACACAAATAA GAAAACGGCA AACCCACCCC ATTTTTTAAT	900
55	TTTATTATTA TTAATTTTT TTGTTGGCAA AAGAATCTCA GGAACGGCCC TGGGCACCTA	960
	CTATATTAAT CATGCTAGTA ACATGAAAAA TGATGGGCTC CTCCTAATAG GAAGGCGAGG	1020
<b></b>	AGAGGAGAAG GCCAGGGGAA TGAATTCAAG AGAGATGTCC ACGGACGAAA CATACGGTGA	1080
60		

	ATAATTCACG	CTCACGTCGT	TCTTCCACAG	TATCTTGTTT	TGATCATTTC	CACTGCACAT	1140
	TTCTCCTCAA	GAAAAGCGAA	AGGACAGACT	GTTGGCTTTG	TGTTTGGAGG	ATAGGAGGGA	1200
5	GAGAGGGAAG	GGGCTGAGGA	AATCTCTGGG	GTAAGAGTAA	AGGCTTCCAG	AAGACATGCT	1260
	GCTATGGTCA	CTGAGGGGTT	AGCTTTATCT	CCICITGITG	ATGCATCCGT	CCAAGTTCAC	1320
10	TGCCTTTATT	TTCCCTCCTC	CCTCTTGTTT	TAGCTGTTAC	ACACACAGTA	ATACCTGAAT	1380
10	ATCCAACGGT	ATAGATCACA	AGGGGGGAT	GTTAAATGTT	AATCTAAAAT	ATAGCTAAAA	1440
	AAAGATTTTG	ACATAAAAGA	GCCTTGATTT	тааааааааа	AGAGAGAGAG	ATGTAATTTA	1500
15	AAAAGTTTAT	ТАТАААТТАА	ATTCAGCAAA	AAAAGATTTG	CTACAAAGTA	TAGAGAAGTA	1560
	ТААААТААА	GTTATTGTTT	GAAAAAAAAA	AAAAAAAAW	CTCGACCGCA	AGGGAAT	161

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# (2) INFORMATION FOR SEQ ID NO: 203:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1974 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 203:

GAATTCGGCA	CGAGGCTGAG	GGAGCTGCAG	CGCAGCAGAG	TATCTGACGG	CGCCAGGITG	60
CGTAGGTGCG	GCACGAGGAG	TTTTCCCGGC	AGCGAGGAGG	TCCTGAGCAG	CATGGCCCGG	120
AGGAGCGCCT	TCCCTGCCGC	CGCGCTCTGG	CTCTGGAGCA	TCCTCCTGTG	CCTGCTGGCA	180
CTGCGGGGGG	AGGCCGGGCC	GCCGCAGGAG	GAGAGCCTGT	ACCTATGGAT	CGATGCTCAC	240
CAGGCAAGAG	TACTCATAGG	ATTTGAAGAA	GATATCCTGA	TTGTTTCAGA	GGGGAAAATG	300
GCACCTTTTA	CACATGATTT	CAGAAAAGCG	CAACAGAGAA	TGCCAGCTAT	TCCTGTCAAT	360
ATCCATTCCA	TGAATTTTAC	CTGGCAAGCT	GCAGGGCAGG	CAGAATACTT	CTATGAATTC	420
CTGTCCTTGC	GCTCCCTGGA	TAAAGGCATC	ATGGCAGATC	CAACCGTCAA	TETCCCTCTG	480
CTGGGAACAG	TGCCTCACAA	GGCATCAGTT	GTTCAAGTTG	GTTTCCCATG	TCTTGGAAAA	540
CAGGATGGGG	TGGCAGCATT	TGAAGTGGAT	GTGATTGTTA	TGAATTCTGA	AGGCAACACC	600
ATTCTCCAAA	CACCTCAAAA	TGCTATCTTC	TTTAAAACAT	GTCAACAAGC	TGAGTGCCCA	660
GGCGGGTGCC	GAAATGGAGG	CTITTGTAAT	GAAAGACGCA	TCTGCGAGTG	TCCTGATGGG	720
TTCCACGGAC	CTCACTGTGA	GAAAGCCCTT	TGTACCCCAC	GATGTATGAA	TGGTGGACTT	780
TGTGTGACTC	CTGGTTTCTC	CATCTGCCCA	CCTGGATTCT	ATGGAGTGAA	CTGTGACAAA	840
GCAAACTGCT	CAACCACCTO	CTTTAATGG	GGGACCTGTT	TCTACCCTGG	AAAATGTATT	900

	TSCCCTCCAG GACTAGAGGG AGAGCAGTGT GAAATCAGCA AATGCCCA	CA ACCCTGTCGA	960
<b></b>	AATGGAGGTA AATGCATTGG TAAAAGCAAA TGTAAGTKTT CCAAAGGT	TA CCAGGGAGAC	1020
5	CTCTGTTCAA AGCCTGTCTG CGAGCCTGGC TGTGGTGCAC ATGGAACC	TG CCATGAACCC	1080
	AACAAATGCC AATGTCAAGA AGGTTGGCAT GGAAGACACT GCAATAAA	AG GTACGAAGCC	1140
10	AGCCTCATAC ATGCCCTGAG GCCAGCAGGC GCCCAGCTCA GGCAGCAC	AC GCCTTCACTT	1200
	AAAAAGGCCG AGGAGCGGCG GGATCCACCT GAATCCAATT ACATCTGG	TG AACTCCGACA	1260
	TCTGAAACGT TTTAAGTTAC ACCAAGTTCA TAGCCTTTGT TAACCTTT	CA TGTGTTGAAT	1320
15	GTTCAAATAA TGTTCATTAC ACTTAAGAAT ACTGGCCTGA ATTTTATT	AG CTTCATTATA	1380
	AATCACTGAG CTGATATTTA CTCTTCCTTT TAAGTTTTCT AAGTACGI	CT GTAGCATGAT	1440
20	GGTATAGATT TTCTTGTTTC AGTGCTTTGG GACAGATTTT ATATTATG	TC AATTGATCAG	1500
	GTTAAAATTT TCAGTGTGTA GTTGGCAGAT ATTTTCAAAA TTACAATC	CA TITATGGTGT	1560
25	CTGGGGGCAG GGGAACATCA GAAAGGTTAA ATTGGGCAAA AATGCGTA	AG TCACAAGAAT	1620
25	TTGGATGGTG CAGTTAATGT TGAAGTTACA GCATTTCAGA TITTATTC	ETC AGATATTTAG	1680
	ATGTTTGTTA CATTTTTAAA AATTGCTCTT AATTTTTAAA CTCTCAA	TAC AATATATTTT	1740
30	GACCTTACCA TTATTCCAGA GATTCAGTAT TAAAAAAAAA AAAATTAG	CAC TGTGGTAGTG	1800
	GCATTTAAAC AATATAATAT ATTCTAAACA CAATGAAATA GGGAATA	TAA TGTATGAACT	1860
35	TTTTGCATTG GCTTGAAGCA ATATAATATA TTGTAAACAA AACACAG	CTC TTACCTAATA	1920
<i>)</i>	AACATTTAT ACTGTTTGTA TGTATAAAAT AAAGGTGCTG CTTTAGT	TTT CTGA	1974

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# (2) INFORMATION FOR SEQ ID NO: 204:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1057 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

#### (vi) SEQUENCE DESCRIPTION: SEO ID NO: 204:

50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 204:					
50	CGGCCTTCCG GGGCAACCGT TCGTCCCAAC NCGGGAAAGG GTCCTGGAGN CGGGAACTAG	60				
	GAGCCTCGGA AGTCCAAGGG CGGAGCGCCC TTTGCTAATA AGCCAATCAG AACGTGAGAC	120				
55	GCTCCGGTGG GNCGGTGCCG TCGAGCGCGG GGTGGACTCT GGGTGACTTG GCTGGCGGGA	180				
	TCAAGTGCAG CTGCTTCAGG CTGAGGTGGC AGATAGTGAG CGCTGGTGGC GGAGTTAAAG	240				
60	TYAAAGCAGG AGAGTAATWA TGAATAGCGC AGCGGGATTC TCACACCTAG ACCGTCGCGA	300				

PCT/US98/11422

	GCGGGTTCTC	AAGTTAGGGG	AGAGTTTCGA	GAAGCAGCCG	CGCTGCGCTT	CCACACTGTG	360
•	CGCTATGACT	TCAAACCTGC	TTCTATTGAC	ACTTCTTCTG	AAGGATACCT	TGAGKTTGGC	420
5	GAAGKTGAAC	AGKTGACCAT	WACTCTGCCM	AATATAGAAA	GTTGAAGGAA	GCAGTAAAAT	480
	TCAGTATCGT	AAAGAACAAC	AGCAACAACA	ATGTGGAATT	CASCCAGGAC	TCCCAATCTT	540
10	GTAAAACATT	CTCCATCTGA	AGATAAGATG	TCCCCAGCAT	CTCCAATAGA	TGATATCGAA	600
10	AGAGAACTGA	AGGCAGAAGC	TAGTCTAATG	GACCAGATGA	GTAGTTGTGA	TAGTTCATCA	660
	GATTCCAAAA	GTTCATCATC	TTCAAGTAGT	GAGGATAGTT	CTAGTGACTC	AGAAGATGAA	720
15	GATTGCAAAT	CCTCTACTTC	TGATACAGGG	NAATTGTGTC	TCAGGACATC	CTACCATGAC	780
	ACAGTACAGG	ATTCCTGATA	TAGATGCCAG	TCATAATAGA	TTTCGAGACA	ACAGTGGCCT	840
20	TCTGATGAAT	ACTTTAAGAA	ATGATTTGCA	GCTGAGTGAA	TCAGGAAGTG	ACAGTGATGA	900
20	CTGAAGAAAT	ATTTAGCTAT	ТАААААТААА	TTATACAGCA	TGTATAATTT	ATTTTGTATT	960
	ААСААТАААА	ATTCCTAAGA	CTGAGGGAAA	TATGTCTTAA	CTTTTGATGA	TAAAAGAAAT	1020
25	TAAATTTGAT	TCAGAAAAAA	. AAAAAAAAA	AACTCGA			105

#### . 30 (2) INFORMATION FOR SEQ ID NO: 205:

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## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 721 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 205:

40	GAATTCGGCA CGAGTCATCC CTCTCCCTCT TTCACTCCCT TACTCTTACT CTGTTTTTTG	60
	TGCTCCAGAC AGACAGACCC TACCTCTTTT GCTTCTTTT TGTTTGTTTG TTTTGAGATG	120
15	GAGTGTCGCT CTTGTTGCCC AGGCTGGAGT GCAGTGGCGC AATCTCGGCT CACCACAACC	180
45	TCTGCCTCCC GGGTTCAAGC AATTCTCCTG CCTCAGCCTC CCGAGAAGCT GGGGATTACA	240
	GGCATGCGCC ACCACACCCA GCTNAATTIT ATATTTTTAG TAGAGATGGT GTTTCTCCAT	300
50	GTTGGTCAGG CTGGCCTCAA ACTCCCAACC TCAGGTGATN CCGCCTGCTT TGGCCTCCCC	360
	AAAGTGCTGG GATTACAGGC GTGAGCCACT GCGCCCAGCC TCTTTTGCTC CTTTATACTC	420
<i>E</i>	ATTAACTCAC GCCTGTAATC CCTGTTTTGG GAGGCCAAAG TGAGAAGGTT GCTTGAGGCC	480
55	AAGAGTTTGA GACTAGCCTG GGCAACACAG CAAGATGCCA TCTTTATAAT AAAAATAAAA	540
	ATAAAAATCA ATTAGCTGGG CATGGTGGAA CGCACCTGTA GTCCCAGCCA ATTGAGAGGC	600
60	TGAAGTGGGA GGATCATTGA GCCCAGGAGT TGAGGTTGCA GTGAGCCATG ATCATGTCAC	660

PCT/US98/11422

•	TACACTCAGC CTGGGCAATA GAGGGACATG TTGTCTCTAA AAAAAAAAAA	720
	A	721
5		
10	(2) INFORMATION FOR SEQ ID NO: 206:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2465 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
15	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 206:	•
20	CCACCATTTA TCCAACTGAA GAGGAGTTAC AGGCAGTTCA GAAAATTGTT TCTATTACTG	60
20	AACGTGCTTT AAAACTCGTT TCAGACAGTT TGTCTGAACA TGAGAAGAAC AAGAACAAAG	120
	AGGGAGATGA TAAGAAAGAG GGAGGTAAAG ACAGAGCTTT GAAAGGAGTT TTGCGAGTGG	180
25	GAGTATTGGC AAAAGGATTA CTTCTCCGAG GAGATAGAAA TGTCAACCTT GTTTTGCTGT	240
	GCTCAGAGAA ACCTTCAAAG ACATTATTAA GCCGTATTGC AGAAAACCTA CCCAAACAGC	300
	TIGCTGTTAT AAGCCCTGAG AAGTATGACA TAAAATGTGC TGTATCTGAA GCGGCAATAA	360
30	TTTTGAATTC ATGTGTGGAA CCCAAAATGC AAGTCACTAT CACACTGACA TCTCCAATTA	420
	TTCGAGAAGA GAACATGAGG GAAGGAGATG TAACCTCGGG TATGGTGAAA GACCCACCGG	480
35	ACGTCTTGGA CAGGCAAAAA TGCCTTGACG CTCTGGCTGC TCTACGCCAC GCTAAGTGGT	540
	TCCAGGCTAG AGCTAATGGT CTGCAGTCCT GTGTGATTAT CATACGCATT CTTCGAGACC	600
	TCTGTCAGCG AGTTCCAACT TGGTCTGATT TTCCAAGCTG GGCTATGGAG TTACTAGTAG	660
40	AGAAAGCAAT CAGCAGTGCT TCTAGCCCTC AGAGCCCTGG GGATGCACTG AGAAGAGTTT	720
	TTGAATGCAT TTCTTCAGGG ATTATTCTTA AAGGTAGTCC TGGACTTCTG GATCCTTGTG	780
45	AAAAGGATCC CTTTGATACC TTGGCAACAA TGACTGACCA GCAGCGTGAA GACATCACAT	840
	CCAGTGCACA GTTTGCATTG AGACTCCTTG CATTCCGCCA GATACACAAA GTTCTAGGCA	900
	TGGATCCATT ACCGCAAATG AGCCAACGTT TTAACATCCA CAACAACAGG AAACGAAGAA	960
50	GAGATAGTGA TGGAGTTGAT GGATTTGAAG CTGAGGGGAA AAAAGACAAA AAAGATTATG	1020
	ATAACITITA AAAAGTGTCT GTAAATCTTC AGTGTTAAAA AAACAGATGC CCATTTGTTG	1080
55	GCTGTTTTC ATTCATAATA ATGTCTACAT TGAAAAATTT ATCAAGAATT TAAAGGATTT	1140
	CATGGAAGAA CCAAGTTTTT CTATGATATT AAAAAATGTA CAGTGTTAGG TATTATTTGA	1200
	ATGGAAAGAC ACCCAAAAAA AAAAATGTGC TCCGACTAGG GGGAAAACAG TAGTTCCGAT	1260
60		

	TTTTTCCCAT	TATTTTTATT	TTATTTTCTG	GTTGCCCTAG	CTTCCCCCCC	TATTTTTGTG	1320
	TCTTTTATTA	ACTAGTGCAT	TGTCTTATTA	AATCTTCACT	GTATTTAATG	CAGGATGTGT	1380
5	GCTTCAGTTG	CTCTGTGTAT	TTTGATATTT	TAATTTAGAG	GITTIGITTG	CTTTTTGACA	1440
	CTAGTTGTAA	GTTACTTTGT	TATAGATGGT	ATCCTTTACC	CCTTCTTAAT	ATTTTACAGC	1500
10	AGTACGTTTT	TTTGTAACGT	GAGACTGCAG	AGTITGTITT	TCTATATGTG	AAGGATTACA	1560
10	ACACAAAAAG	TTATCCTGCC	ATTCGAGTGC	TCAGAACTGA	ATGTTTCTGC	AGATCTTGTG	1620
	GCATTTGTCT	CTAGTGTGAT	ATATAAAGGT	GTAATTAAGA	CAGAGTTCTG	TTAATCTAAT	1680
15	CAAGTTTGCT	GTTAGTTGTG	CATTAGCAGT	ATAAAAGCTA	ATATATACTA	TATGGTCTTG	1740
	CAACAGTTTT	AAAGCCTCTG	CATAATTGAT	AATAAAAATG	CATGACATTC	TTGTTTTTAA	1800
20	TAGACTTTTA	AAATCATAAT	TTTAGGTTTA	ACACGTAGAT	CTTTGTACAG	TTGACTTTTT	1860
20	GACATAGCAA	GGCCAAAAAT	AACTTTCTGA	ATATTTTTT	CITGTGTATA	AGTGGAAAGG	1920
	GCATTTTTCA	CATATAAGTG	GGCTAACCAA	TATTTTCAAA	AGAACTTCAT	CATTGTACAA	1980
25	CTAACAACAG	TAACTAGCCC	TTAATTATGG	TGACAGTTCC	TTATTGGTGT	GTGTGAGATT	2040
25	ACTCTAGCAA	CTATTACAGT	ATAACACAGA	TGATCTTCTC	CACACACCCC	ATCACCCAGA	2100
30	TAATTTACAG	TTCTGTTAAC	AGTGAGGTTG	ATAAAGTATT	ACTGATAAAA	AATTATCTAA	2160
50	GGAAAAAAAC	AGAAAATTAT	TIGGIGIGG	CATCTTACCT	GCTTATGTCT	CCTACACAAA	2220
	GCTAAATATT	CTAGCAGTGA	TGTAATGAAA	AATTACATCI	TACTGTTGAT	ATATGTATGC	2280
35	TCTGGTACAC	AGATGTCATT	TTGTTGTCAC	AGCACTACAG	TGAAATACAC	AAAAAATGAA	2340
	ATTCATATAA	TGACTTAAAT	GTATTATATO	TTAGAATTGA	CAACATAAAC	TACTTTTGCT	2400
40	TTGAAATGAT	GTATGCTTC	GTAAAATCAT	ATTCAAATT	KAAAAAAAA 1	AAAAAAAAA	2460
<b>-7</b> 0	CTCGA						2465

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## (2) INFORMATION FOR SEQ ID NO: 207:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1480 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 207:

GAATTCGGCA CGAGCTCAAG CTGGCAGGTG GTCGGGGGAG CGGCCGGAGA GGAGCTGCCG 60
GGAGTTCGTG CCCTGCAGGA CATGACACCA GTGGCATATC ACGGCCATGG GGTCTCAGCA 120

TTCCGCTGCT GCTCGCCCCT CCTCCTGCAG GCGAAAGCAA GAAGATGACA GGGACGGTTT 180

	GCTGGCTGAA	CGAGAGCAGG	AAGAAGCCAT	TGCTCAGTTC	CCATATGTGG	AATTCACCGG	240
_	GAGAGATAGC	ATCACCTGTC	TCACGTGCCA	GGGGACAGGC	TACATTCCAA	CAGAGCAAGT	300
5	AAATGAGTTG	GTGGCTTTGA	TCCCACACAG	TGATCAGAGA	TTGCGCCCTC	AGCGAACTAA	360
	GCAATATGTC	CTCCTGTCCA	TCCTGCTTTG	TCTCCTGGCA	TCTGGTTTGG	TGGTTTTCTT	420
0	CCTGTTTCCG	CATTCAGTCC	TTGTGGATGA	TGACGGCATC	AAAGTGGTGA	AAGTCACATT	480
	TAATAAGCAA	GACTCCCTTG	TAATTCTCAC	CATCATGGCC	ACCCTGAAAA	TCAGGAACTC	540
	CAACTTCTAC	ACGGTGGCAG	TGACCAGCCT	GTCCAGCCAG	ATTCAGTACA	TGAACACAGT	600
15	GGTGAATTTT	ACCGGGAAGG	CCGAGATGGG	AGGACCGTTT	TCCTATGTGT	ACTTCTTCTG	660
	CACGGTACCT	GAGATCCTGG	TGCACAACAT	AGTGATCTTC	ATGCGAACTT	CAGTGAAGAT	720
20	TTCATACATT	GGCCTCATGA	CCCAGAGCTC	CTTGGAGACA	CATCACTATG	TGGATTGTGG	780
	AGGAAATTCC	ACAGCTATTT	AACAACTGCT	ATTGGTTCTT	CCACACAGCG	CCTGTAGAAG	840
25	AGAGCACAGC	ATATGTTCCC	AAGGCCTGAG	TTCTGGACCT	ACCCCACGT	GGTGTAAGCA	900
25	GAGGAGGAAT	TGGTTCACTT	AACTCCCAGC	AAACATCCTC	CTGCCACTTA	GGAGGAAACA	960
	CCTCCCTATG	GTACCATTTA	TGTTTCTCAG	; AACCAGCAGA	ATCAGTGCCT	AGCCTGTGCC	1020
30	CAGCAAATAG	TTGGCACTCA	ATAAAGATTI	GCAGAATTTA	ATACAGATCT	TTTCAGCTGT	1080
	TCTTAGGGCA	A TTATAAATGG	AAATCATAAC	GTGGTTCTAG	GTTATCAAAC	CATGGAGTGA	1140
25	TGTGGAGCT	A GGATTGTGAG	TGACCTGCAC	GCCATTATC	GTGCCTCATC	TGTGCAGAAG	1200
35	TCGCAGCAG	A GAGGGACCAT	CCAAATACCT	r aagagaaaa	AGACCTAGTC	AGGATATGAA	1260
	TTTGTTTCAC	G CTGTTCCCA	AGGCCTGGG	A GCTTTTTGA	A AAGAAAGAA?	AAAGTGTGTT	1320
40	GGCTTTTTT	r ttttttaga/	AGTTAGAAT	r GTTTTTACC	A AGAGTCTATO	TGGGGCTTGA	1380
	TTCACCCTT	C ATCCATTGGG	TGGAACATG	G ATTGGGGAT	r tgatagaaa	A ATAAACCCTG	1440
15	CTTTTGATT	C AAAAAAAAA	AAWAAAAA A	A AAAAACTCG	A		1480

# (2) INFORMATION FOR SEQ ID NO: 208:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 872 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

55 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 208:

CAGTATTTCC CTCAGTACTG TAAGCAAAAG TGGTATGTTT TTCTTTCTTT ATGTCTACTC

600

60

	TGTCCTCTGT GGCCTTCTGG TGTACCCCTC TCTTCCTAGC CATTCAGTCT CTCTAGTCAC	120
	CTCCCTAGTA GCTAGTGCTC TCTAAGTTTT TATTTAATTA GAACAACTCC ATTTCCATTT	180
5	CAAGGTAGGT CAATGGGGGG AAAAGCCTCA TGATTTAAAC TGAAGTTAAC AACACAGCTT	240
•	TTAAAATGAA AACTCATACT CCAACTTCTA AAGTATATTT GAGCTGATTT GTTTCCAAAA	300
	CAAAGATATG CTGTACCTAA AACTGCTAAA ACAAAAATAT AAAGACAAGG ACTAGGTGAT	360
10	TAAGGGGAGA GAAAAATCAT YTCTTTTCCA GGAAACCTTT GCTAAAATAA GCAAAACTTG	420
	ANTICTATGCT TCATGGAAAC TGACACAAAG AAAAGAAACT GATGGATTGC ACAGGCCTTG	480
15	TTATAGAAAT AGATCTATAA AAAGATCTGT CCACAGGAAA TATACACCTT CTCCTGGTTC	540
	TGAACTTCAA TGGGGATTTG TCACCTAGGT CTCCATCTAT AGGAATACCT TCACATACCT	600
20	ATCTATTCAT GCACATATTC TGAAAACAGG TACATACAAA ATTACAACAA AGGAAAAAAA	660
20	TTCTATTGAA CACTTAAAAA TAGAAACAGG CCAGGCACGG TGGCTCATGC TGTAATCCCA	720
	ACAATTTGGG AGGCTGAGGC TGGTGGATCA CCTGAGGTCA GGAGTGTGAG ACCAGCTTGG	780
25	CCAACATGGT GAAACCCCGT CACTACIAAA AATACAAAAA AAATTAGCCT GTGTGGTGGC	840
	ACACTCMTAC AATCCNGGCT GACTCGGGAA AN	872
30		
50	, , , , , , , , , , , , , , , , , , ,	
	(2) INFORMATION FOR SEQ ID NO: 209:	
35	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1779 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 209:	
	AATTGCCAAG ACTGCACAAA ATTACAGTGC TAATGTATAT GGTTGCAGTT CACATAAAGA	60
A 5	CAAAAGCATC TGTTATGAAA TGAGTAGTAA TATTGGGTGG TTGATTTGTT CTTAGCAGAC	120
45	TTGGCTTCAT WTTGGTCTTG AGATAAAATG GCCAGCATAA ATGCTGTTTA TATTCACGTT	180
	TTCCTAGGTG TGTGTGCA GGCCACAGCA GCATGCCCTT GGTGTAGTCA GTGCCGAAAS	24
50	GGGTCTGTTC CTTCTTGAGC CTGCCTGCAG GGATGGTCTC CTTTTAAAGC AGGTTGTGTG	30
	CAGCATTCAG TACACTGAAG GTAAGCTAAA CCATCAACAT CTCTGGTGTT TTAAGATGTT	36
55	ATTITATIOG AACAACIGAC AAAIGAGGGA TGITAGCITT GIGGCAGAAT ICCCIGCAIG	42
در	TGTGATAACT GATCTTGTTT TATTTTTTGG CATTGCAACT GTGGCATAGT TACAATTTCT	48

GTTTGKTCAT CACATTTAAA ATTGGRAGAG AACGCGCTTG AKGGATAGAG CGCCTTCAGK

GTACTGTTTC TTATTAACTT TACTTTTTTT AAATCAACTT GCTATAGACT TTATATACAT

	TTTGTTAAAT A	TAGTTCCTA	GTGACATAGA	AACGATGCGT	AGTTTTCATT	TACTAATTAC	660
_	AAATGTTGAG G	CCTAATTCT	GAAAGTCCTC	ATATTTAAAG	GCTAGACAAC	GTAATGAAAT	720
5	TTTTAACTAT T	TGTATGTCA	TTTTGAAAGT	GTACTGCTTT	ATGGTAAAAG	TGTTTTTCAT	780
	TIGITCATIG T	TTTCATTAT	TTGTGATCAT	GTTGTCTTTC	AATACAGGCA	TAAACCTTCC	840
0	ACTCTTGAAC A	AAGCAGCTG	CTTTTTAAAA	GCGGTAATTG	CTTCTTTACC	TTTTATTTCT	900
	TTTGTAAATG A	AGCTTTTCT	TTAAGAATGT	GACTTTAAAG	TGTTGTCTAT	TGCATAAAAC	960
15	AGTTGACACT C	CACTTATTGT	AAAGTGAAGA	TIGTICTACT	GCATGTGAAG	TGGACCATGC	1020
13	AGATTTCTGT A	ATGITCTCAG	TATGCATCAC	TAGATAATAA	AGTCTTTTGT	GAACAAGGCA	1080
	TTTGTAGCCA T	TTTTTAAAAG	TTTTTGTCTT	CAGTGCTGGT	AAGTCAGGTA	AACCATAAAT	1140
20	AGTTAAAAGC A	ACCTTTIGT	TTTTTTCCTG	AAAGTTTTTA	ATTGAAAGTA	TTATTAGTTA	1200
	AAGATGTAAA (	CTAGCCAAA	ATTACCAGTT	TATTAATAAT	TAGGATCCTA	ATTATTTCAA	1260
25	AAAATCCTAC A	AAATATTGTC	AGCTTTCAGT	GTAGTGAGAT	TATTCCTGTA	GCTTATCCCC	1320
25	TATAATTCAG (	GATTTAACTA	ATGTTTCTGC	TATITICICA	CTTTTCCTTT	TGATGGTGCG	1380
	GAAAGAGAAA	AAGGAAAACG	GGGCACAGGC	: CATTCGACGO	CTTCTCCAAG	GGGTCTGATT	1440
30	TGCTGAGACA	CCAGCTTCAC	CTTCTTAACA	AGGCACCTA	TTACAACAAC	CATGCACATT	1500
	TTGGTGCATT	CAAGAATGGA	AAATCAGAAT	AGCAGCATTO	ATTCTTCTGG	TGCAGCTCAG	<sub>.</sub> 1560
25	TGGAAGATGA	TGACAACCAG	AAGACATGAG	CTAAGGGTA	A GGGACTGTTC	TGAAGAACCT	1620
35	TTCCATTTAG	TGATCAAGAT	ATGGAAGCT	ATTTCTGAA!	A ATGCTCAGTO	TGTACTCTAA	1680
	TTATTTATGG	TACCATTTGA	ATTGTAACT	r GCATTTTAGG	C AGTGCATGT	TCTAATTGAC	174
40	TTACTGGGAA	ACTGAATAAA	ATATGCCTC	TATTATCAA			177

#### 45 (2) INFORMATION FOR SEQ ID NO: 210:

50

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2110 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 210:

	GCCTCGGAGG GGCCTCGGCT GCCCCACCCT CGGAGCCACT GCTAGAAGGG GCCGCTCCCC	240
	AGCCTTTCAC CACCTCTGAT GACACCCCCT GCCAGGAGCA GCCCAAGGAA GTCCTTAAGG	300
5	CTCCCAGCAC CTCGGGCCTT CAGCAGGTGG CCTTTMAGCC TGGGCAGAAG GTTTATGTGT	360
	GGTACGGGGG TCAAGAGTGC ACAGGACTGG TGGWGCAGCA CAGCTGGATG GAGGGTCAGG	420
10	TGACCGTCTG GCTGCTGGAG CAGAAGCTGC AGGTCTGCTG CAGGGTGGAG GAGGTGTGGC	480
10	TGGCAGAGCT GCAGGGCCCC TGTCCCCAGG CACCACCCCT GGAGCCCGGA GCCCAGGCCC	540
	TGGCCTACAG GCCCGTCTCC AGGAACATCG ATGTCCCAAA GAGGAAGTCG GACGCATGGA	600
15	AATGGATGAG ATGATGGCGG CCATGGTGCT GACGTCCCTG TCCTGCAGCC CTGTTGTACA	660
	GAGTCCTCCC GGGACCGAGG CCAACTTCTC TGCTTCCCGT GCGGCCTGCG ACCCATGGAA	720
20	GGAGAGTGGT GACATCTCGG ACAGCGGCAN CAGCACTACC AGCGGTCACT GGAGTGGGAG	780
20	CAGTGGTGTC TCCACCCCCT CGCCCCCCCA CCCCCAGGCC AGCCCCAAGT ATTTGGGGGA	840
	TGCTTTTGGT TCTCCCCAAA CTGATCATGG CTTTGAGACC GATCCTGACC CTTTCCTGCT	900
25	GGACGAACCA GCTCCACGAA AAAGAAAGAA CTCTGTGAAG GTGATGTACA AGTGCCTGTG	960
	GCCAAACTGT GGCAAAGTTC TGCGCTCCAT TGTGGGCATC AAACGACACG TCAAAGCCCT	1020
30	CCATCTGGGG GACACAGTGG ACTCTGATCA GTTCAAGCGG GAGGAGGATT TCTACTACAC	1080
30	AGAGGTGCAG CTGAAGGAGG AATCTGCTGC TGCTGCTGCT GCTGCTGCCG CAGACCCCCA	1140
	GTCCCTGGGA CTCCCACCTC CGAGCCAGCT CCCACCCCCA GCATGACTGG CCTGCCTCTG	1200
35	TETGETETTE CACCACCTET GCACAAAGCC CAGTCCTCCG GCCCAGAACA TCCTGGCCCG	1260
	GAGTCCTCCC TGCCCTCAGG GGCTCTCAGC AAGTCAGCTC CTGGGTCCTT CTGGCACATT	1320
40	CAGGCAGATC ATGCATACCA GGCTCTGCCA TCCTTCCAGA TCCCAGTCTC ACCACACATC	1380
.0	TACACCAGTG TCAGCTGGGC TGCTGCCCCC TCCGCCGCCT GCTCTCTMTC TCCGGTCCGG	1440
	AGCCGGTCGC TAAGCTTCAG CGAAGCCCCA GCAGCCAGCA CCTGCGATGA AATCTCATCT	1500
45	GATCGTCACT TCTCCACCCC GGGCCCAGAG TGGTGCCAGG AAAGCCCCGAG GGGAGGCTAA	1560
	GAAGTGCCGC AAGTGTATGG CATCGAGCAC CGGGACCAGT GGTGCACGGC CTGCCGGTGG	1620
50	AAGAAGGCCT GCCAGCGCTT TCTGGACTGA GCTGTGCTGC AGGTTCTACT CTGTTCCTGG	1680
	CCCTGCCGGC AGCCACTGAC AAGAGGCCAG TGTGTCACCA GCCCTCAGCA GAAACCGAAA	1740
	GAGAAAGAAC GGAAACACGG AGTTTGGGCT CTGTTGGCTA AGGTGTAACA CTTAAAGCAA	1800
55	TTTTCTCCCA TTGTGCGAAC ATTTTATTT TTAAAAAAA GAAACAAAAA TATTTTTCCC	186
	CCTAAAATAG GAGAGAGCCA AAACTGACCA AGGCTATTCA GCAGTGAACC AGTGACCAAA	192
60	GAATTAATTA CCCTCCGTTT CCCACATCCC CACTCTCTAG GGGATTAGCT TGTGCGTGTC	198
~ ~		

	AAAAGAAGGA	ACAGCTCGTT	CTGCTTCCTG	CTGAGTCGGT	GAATTCTTTG	CTTTCTAAAC	2040
	TCTTCCAGAA	AGGACTGTGA	GCAAGATGAA	TTTACTTTTC	TTAAAAAAA	АААААААА	2100
;	AAAAACTCGA	•					2110

10 (2) INFORMATION FOR SEQ ID NO: 211:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 938 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 211:

20	GGCACAGGAA AAAAAAGAAA AAAGAAAAAA GAAAAAAGTT TTTGTACCCA CAGATTAGCA	60
	TTTTCTTGAT GTTTGAAAAA AGTTTAAGCT ATGTCCTAAT TTAAAAATGA GCACAAACTA	120
25	CTTAACAGAT GTCTGTTCCC TCTTCTCTTA CTTAAATTAT CTTTATTTTC ACCATCACCT	180
25	CCCAGUGCCG AACACCTGAN CTCTGTGTTT TGTGGTTGGA TCCTGGGTTG CCAAGTTCCT	240
	ATTTGGTCAG TCCCTGGCCT GTGGGGGGGT CTCAGGAAGT GGCATGCTCT TCAMGRAGGA	300
30	TOGTICATYT CCAGTATAAC CAWITTGITA ATAATAGITG ATAATTCCCA GCTTTTACCA	360
	GATGARTTTT GACTTATTTT TCCTCCTTTG ACCTGTTCAA AGCTAACATA TCTCGGTCAG	420
	TTCGGAGAGG GTGGGGGATT TGAGAATGTG AGGAGGAGTG GGGTTAGAAT GGGTTTGCCT	480
35	ATCTGGGCAA GGAAAGAGTT CCTAGTCGAT TGGGCACAAT GACAAAATGA TTCCATGGAT	540
	AGAATCGTCC CATGTTGCTG GAACACCTCA CGTGTTGTGA ACGCCTTAAA TTCCTGCCAT	600
40	CCCTTCTCTG ATTCCCCACC TCCCTGTAGT TTCCACAGGA TTTATCTCTC TGTACCCCCG	660
	TCCTCCAACT CTACTCTGTC AGCCTCTCCT CCATCCCTTA CTTCCCTTCT AAATTCCAGG	720
	AGATGACCIC ACTITIGCAAA GCAAATTGGA GCCACCAAAT TGTAGCTCTC CTCGGTGGAA	780
45	ACTGCATCTG TGCTCATCCC TGCACCTTCT TGCAGAAAGC CGCCCCCTCA GGCCAAGATG	840
	AGTGCCTGGC CCCCATGGGA GACTCAGACA CTTTGACCCC TTGTGACTTC AGCATCTCCC	900
50	TCTTTAAAGA TTCTCTCCCA ACATTCAGTC GTGCTCGA	938

55 (2) INFORMATION FOR SEQ ID NO: 212:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1551 base pairs

(B) TYPE: nucleic acid

60 (C) STRANDEDNESS: double

# (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 212:

	· · · · -	
5	AGGCTGGACT AAGCATAGAG AACCAGGAGA GAAAGAAAGA TTTAAGAGAC TGAGTAATAT	50
	TTTTTGACAG ATCATTTAAG AAACTGAGTA ATTTTTTTTT TCTCCAAAAG GGCATGGGTT	120
10	TTTTTTTTGT TTTGTTTTTT CTCTATTTGG CACTTTCTAG GGATTGGTCT ATAAATTTTT	130
	TGAAAGATCA TAGGATAAAT TTCTTTGTAG CAACTTCCTA TTTTAGTGTT TATGTTAGGG	240
	GARCCCCARG TGTCCCTGCT GATACGCCAT TAGGGCCACT TCTCAGCCTC TGGCTACATC	300
15	ATAATGCTTT TTTTTCTATC TIGCCAAAGT TTCCMGAAAA TTKAKGTTTT CTAATTTTAA	350
	AAAAATTGGT TGTGGAGATG GGATGGGACC TCTTTATAAG CCCTGAAAAT AAGTGATTTN	420
	TTTTAAGTGC TATTCTGCTA TAAACCTGAT TCTCACTTTT TTCTGTAGAC AACAGTTTTT	430
20	TATAATATAT CTATTTTGTG TGGACATTAT TTCCTTTTAA CCAATACTGA AATTCCATAG	540
	TGTAWACTTT CTCCACATTT TCTTTGATTA ATACTTYCTT AAAATAGACA CTTCGATTCG	600
25	CACCAGCTGT CACCAATAAA GCTGCCCTGA ACATTGTCAA TCAATCCTGT TAACCAATTT	650
	GAGAATTTTT CTGGAATGCT TAGTTAGGGA TGAAATTGCT GGGTTATAGG TATGAGTATG	720
20	CTTGATATAC TTTTCTCCAG AATGTCTACA CCTGTGTGTA CACCACATCT CCAGAGATAG	730
30	GGGAATCITA TGTCCCTGCT AACTGCTCTC GTTATTTAAT TTTCTGACAT TTGCCGCCGC	840
	CGCCGCCCCC TGCCCCCAAC ACACACATGG TATAAAGTGG TAGTTTCTTG TTTTAAATTG	900
35	AACTTTGAA TGATTTGAAT TTGGGCATTT CTTTGTATCC TGAGTTATTT TGGTTTCCCG	960
	TTATGTGAAT ATCCTTTTCC TATGCTTTAA CTACTTTTCT AATTTGTCCC TTTTTTTNGGT	1020
40	TATCAAATTC CAGGCCATTG TCTATTCCAT CGTCACTTTT GGGTATTGGA AACATCTTTC	1030
40	CATTCTGTAG CCTGTCTGTT GAACATAAAT CTTGATTTTT ATGTAATCAG ATTTTTCTCC	1140
	TTACGGTTAT GTTCTTGGAA TTTTATTTAA GAAATCTTTT TCTATCCTGA GACCACAAAA	1200
45	ATGTCCCCAC CATTTTCTTC TGTTTCATAG TTTTGCCTTG TATGTTTAAT CCTTTAAGGC	1250
	ATGTGTAGTT CATTTTATAT GGTGTGAAAT AGTTCTTATT CATTTATTCA ACACATATTG	1320
50	GTGGAGTGCC TGCTGATGGT AGTACTCTTC AGAGTACTTT GTATATATTT GTGAACACAT	1380
50	ATTCTTGCCC TGGAAGCTTA TGTTGTCNTT CAAGGTAGAT CCNTACTCGG TTTCCACCTG	1440
	THITCTICAG CCCTCAGGAT GAATTCCACA ATTITACACA TAGCACCAGT TAAGGAATAG	1500
55	GCTTTATTGG AGAAAAGGAA GGCTTATTAG ACCAGCATCA GCAAAAAAAA A	1551

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 997 pase pairs(B) TYPE: nucleic acid

5	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NG: 213:	
0	AGAGAGTCCT CAACAGAACC TAATCATGCT GGCACCCTAA TOTCATACTT CTAGCCTCCA	50
	GAACTGAGAG AACATAAACT CCAGTTGTTT AAGCTACCCA GECTAIGGIA ITTGTTAITA	120
15	TAGCCCAAGC TAAGTCAGGT GGAAAGGCAG AAATATTTTG AGAAGARTCA TTTCTACAAA	130
IJ	AACAGAGTTG TTCTAAATGA AATGGCCAGA TATTTCATCT TCTTCATACT AGTATTTAIG	240
	AAAGTTTCAT TAAACACCAC TTGGCCAGCA CCCAGGCCTG CCACCCTCAG AACGGCAAAC	300
20	AAAAGCAAAT GATTTGAGGA ACAAAAGAGT GGACACAGAG CTTCTCAGAA GATGGCTCCA	360
	TCTTCTGAGA TGATCTTCTG AGATCATCAA TTTTCTGCAC CTGATGTCCT ACTCCAATTG	420
25	TAGTAGATAA GAGCAAAGAC ACTTCCTGAT CCTGTGGAAA ATGCTGGAGC CCTGCTGATG	430
23	GAGAGGCTGA CACTGGGACC AACAGAAGGC CGGACATTTA TUTGUTGCAG CCCTTCTGCA	540
	CCTGGGCCCT CTTCAGGCCT TGTACCTTGC ACTCCCCATG CCACTGTAGC ACCTGGTAAG	600
30 <sup>°</sup>	CTGAAGTTAG GTATTTGAAG AGATAATTTG CCCCCAACAA AGAATTACTT AAAAGAAAAA	550
	GGAAACCACT AAATTCCACT TGACAAACCA GTTTGTTCAG TITTGACTIT TGCAAATTTG	720
35	AAACTITCTC TITGGCACCA TATGATTCTG TTACATTAGG GCTCATCAAT GCTAAGATAC	780
<i>J J</i>	ACAGCTAGGT CTACCAGCTG CCAGTGGTCA AGAATGAAAG AACCTCTCAG AGAGAGATTA	84
	GTTTCTAATA ACCTAACAGT TTTCCTTGGS TATTACMAAA AAAAAAAAA TTAGAATAAA	90
40	ATGTCAGTGC CATGCAGGCA AGTACAGATA TGGAAATGAA AGCTCTGTCT ACAACTGCAA	96
	GATTTGTTTG TTAATAAAAT TGATTGGGAT CACTCGA	99
45		
	(2) INFORMATION FOR SEQ ID NO: 214:	
	(i) SEQUENCE CHARACTERISTICS:	
50	(A) LENGTH: 1496 base pairs (B) TYPE: nucleic acid	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 214:	٠
	GAATTCGGCA CGAGTGACCA CAGATATCTT TGGCTTTCAG CCTCACCACA ATGCTGTCCA	6
	CTATGTTTT TTTAATCGAT TGACATCTCA TGAATCCACA AATTTAGCCG CTTTTCCATC	12
60	CIRIOTITI IIIANICONI ISSANICONI ISSANICONI	

•	TTTTCCATCT TTGTCATAGC TTCATCACGC ACGATGGAGG TCACTTCAGC ACTATCCGGA	180
	GCGGCCTCAC GGACAGATCR GTGAATTTCC TTTTCCTTTT TCTTGATGTA CCGGATTGTC	240
5	GACTCGTTAA CATTGAGCTC ATGGCCAACA GCACTGTAAC TCATGCCTGA TTGGAGCTTA	300
	TCCAACACGC GGAMTTTCTC CGTAAGGSAM ATCAMGGTCT TCTTTCGCTT AGGAACACTG	360
	GGCARARCTT AARCACTACG CTTGGGGGCC ATTTTAGAAA GCAAAACCAC CCACAAAAAG	- 420
10	CAGAAAAAAA AGTGTCAGTA AACAGACTGN NGANAGGACT CTTTGTTTAC AGCACAGGAG	480
	CTGCGACTAG AAGGCGGCGC TTCTCCCCAG TTCAAACTTC AGCTGGGAAC CTTACCTCCG	540
15 .	CCAACTCCAA ATTTTCACCC TCTGCGCATG CCCGGGAAAS AAACCCCCAG AACAGTACCG	600
	TGATGATTGA TTTTAGGGTT ACAAATACAT TTTAGCAAGT AAGTGAATTT GGCATTACGA	660
20	ATTAATGATT AATGAAGGTC ACCTGTATTT CCATAGATAT GTAATTTTAT TTAAGCAGGT	720
20	TTATTATATT AAGGCGGGGA GGCAGCGCCG AAGACTACAA GTTCCAGCAT GCACCGCGTC	780
	CGGGCGGGTT CGGGCTCCCA GCGAGGGCTT CAGGGACGCC AGCCCGGAGG CATCGGCCGG	840
25	AAGTGTCGTA GGGCAACCAC GTAGTACTCT CTGCGCATGT GCAAAGCGCT GTCGGGGGCC	900
	GCCCTAGCTG CCGTCGCCGC CGCCGGGGCT CTATGGTCTC TCCCTAGAGC TTTGCCGTTG	960
30	GAGGCGGCTG CTGCGGTCTT GTGAGTTTGA CCAGCGTCGA GCGGCAGCAA CATGGAGGAA	1020
50	TTCGACTCCG AAGACTTCTC TACGTCGGAG GAGGACGAGG ACTACGTGCC GTCGGGTGAG	1080
	CGATTCCGCC TGAGGCGAGA AGCGAATTGC CCCGCCCCAC GCCTCACGTG AGGCGCGCTC	1140
35	TGCCCCCGCG GGCGTCTGCC CTGTGGCCCA GGTGGTCCAG GGGGGCTCCT GTTCTCGAGC	1200
	GTCCGCTCCC TCAGGCCCCT CATQCTCGGC CGCTCCGGCC CGAGGCGTGT GCGCGTGGCG	1260
40	GTTCTGTGCT CCCCTCCCGT TGGGCAGCTC CGGCCGCCGC CCCCTCTTGC AGCGCGGGAA	132
70	CGGCACATGG ACACGGCCCC TTGTCGCTAG GGACGCTCGT CGGTCAGCCC CGAACGACAA	138
	CGCTGCTTCA GAAGTCGGGG CGGCAGTTCG AGCCTTGGAA GTTTTTTTCA GCCCTGGCCC	144
45	GAGAGAGCTG CTGGCCAACA ACCCGTCCAA GATAGAGCTG TCCGNTCTCC GNCTGG	149

### 50 (2) INFORMATION FOR SEQ ID NO: 215:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1308 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 215:

	CTGCCTTTGA CCCATCACAC CCCATTTCCT CCTCTTTCCC TCTCCCCGCT GCCAAAAAAA	120
5	AAAAAAAAGG AAACGTTTAT CATGAATCAA CAGGGTTTCA GTCCTTATCA AAGAGAGATG	180
J	TGGAAAGAGC TAAAGAAACC ACCCTTTGTT CCCAACTCCA CTTTACCCAT ATTTTATGCA	240
	ACACAAACAC TGTCCTTTTG GGTCCCTTTC TTACAGATGG ACCTCTTGAG AAGAATTATC	300
0	GTATTCCACG TTTTTAGCCC TCAGGTTACC AAGATAAATA TATGTATATA TAACCTTTAT	360
	TATTGCTATA TCTTTGTGGA TAATACATTC AGGTGGTGCT GGGTGATTTA TTATAATCTG	420
	AACCTAGGTA TATCCTTTGG TCTTCCACAG TCATGTTGAG GTGGGCTCCC TGGTATGGTA	480
15	AAAAGCCAGG TATAATGTAA CTTCACCCCA GCCTTTGTAC TAAGCTCTTG ATAGTGGATA	540
	TACTCTTTTA AGTTTAGCCC CAATATAGGG TAATGGAAAT TTCCTGCCCT CTGGGTTCCC	600
20	CATTTTTACT ATTAAGAAGA CCAGTGATAA TTTAATAATG CCACCAACTC TGGCTTAGTT	660
	AAGTGAGAGT GTGAACTGTG TGGCAAGAGA GCCTCACACC TCACTAGGTG CAGAGAGCCC	720
25	AGGCCTTATG TTAAAATCAT GCACTTGAAA AGCAAACCTT AATCTGCAAA GACAGCAGCA	780
23	AGCATTATAC GGTCATCTTG AATGATCCCT TTGAAATTTT TTTTTTGTTT GTTTGTTTAA	840
	ATCAAGCCTG AGGCTGGTGA ACAGTAGCTA CACACCCATA TTGTGTGTTC TGTGAATGCT	900
30	AGCTCTCTTG AATTTGGATA TTGGTTATTT TTTATAGAGT GTAAACCAAG TTTTATATTC	960
	TGCAATGCGA ACAGGTACCT ATCTGTTTCT AAATAAAACT GTTTACATTC ATTATGGGGT	1020
35	ATGTATGACC TTCATTTTCC AAGAAATAGA ACTCTAGCTT AGAATTATGG ATGCTCTAAA	1080
<i>55</i>	ATGTCAGAAT GGGAACTCTC CTCGAAGTTC TCCCAAACTC AGAGACAGCA CTGCCTTCTC	1140
	CTAAATGATT ATTCTTTTCT CCCTGTTTTC TGGTATTTTC TAGGCATCCT TCTCACCACA	1200
40	GCCATAACCC TTTTTTACTT CCATTAGGCC GTATAACTGG NGGGACNGCT GGTCGGTATA	1260
	TAATACTGGT WCCAACAMAG GGGTTCTGGA TGTACACMAG GTTATCTT	1308
45		
7.7	(2) INFORMATION FOR SEQ ID NO: 216:	
	(i) SEQUENCE CHARACTERISTICS:	
50	(A) LENGTH: 1705 base pairs	
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(D) TOPOLOGY: linear	
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 216:	
	TYCCCATCGA ACCCCTAGAA CGTTTAGATT TTGAAACAGC AAAGAAGGAT TTCCTTGGAT	60

CTGGAGACCC CAAAGAAACA AAGATGCTAA TCACCAAACA GGCTGACTGG GCCAGAAATA

	TCAAGGAGCC CAAAGCCGCC GTGGAGATGT ACATCTCAGC AGGAGAGCAC GTCAAGGCCA	180
	TCGAGATCTG TGGTGACCAT GGCTGGGTTG ACATGTTGAT CGACATCGCC CGCAAACTGG	240
5	ACAAGGCTGA GCGCGAGCCC CTGCTGCTGT GCGCTACCTA CCTCAAGAAG CTGGACAGCC	300
	CTGGCTATGC TGCTGAGACC TACCTGAAGA TGGGTGACCT CAAGTCCCTG GTGCAGCTGC	360
10	AGTGGAGACC CAGCGCTGGG ATGAGGCCTT TGCTTTGGGT GAGAAGCATC CTGAGTTTAA	. 420
10	GGATGACATC TACATGCCGT ATGCTCAGTG GCTAGCAGAG AACGATCGCT TTGAGGAAGC	480
	CCAGAAAGCG TTCCACAAGG CTGGGCGACA GAGAGAAGCG GTCCAGGTGC TGGAGCAGCT	540
15	CACAAACAAT GCCGTGGCGG AGAGCAGGTT TAATGATGCT GCCTATTATT ACTGGATGCT	600
	GTCCATGCAG TGCCTCGATA TAGCTCAAGA TCCTGCCCAG AAGGACACAA TGCTTGGCAA	660
20	GTTCTACCAC TTCCAGCGTT TGGCAGAGCT GTACCATGGT TACCATGCCA TCCATCGCCA	720
20	CACGGAAGAT CCGTTCAGTG TCCATCGTCC TGAAACTCTT TTCAACATCT CCAGGTTCCT	780
	GCTGCACAGC CTGCCCAAGG ACACCCCCTC GGGCATCTCT AAAGTGAAAA TACTCTTCAC	840
25	CTTGGCCAAG CAGAGCAAGG CCCTCGGTGC CTACAGGCTG GCCCGGCACG CCTATGACAA	900
	GCTGCGTGGC CTGTACATCC CTGCCAGATT CCAAAAGTCC ATTGAGCTGG GTACCCTGAC	960
30	CATCCGCGCC AAGCCCTTCC ACGACAGTGA GGAGTTGGTG CCCTTGTGCT ACCGCTGCTC	1020
30	CACCAACAAC CCGCTGCTCA ACAACCTGGG CAACGTCTGC ATCAACTGCC GCCAGCCCTT	1080
٠.	CATCTTCTCC GCCTCTTCCT ACGACGTGCT ACACCTGGTT GAGTTCTACC TGGAGGAAGG	1140
35	GATCACTGAT GAAGAAGCCA TCTCCCTCAT CGACCTGGAG GTGCTGAGAC CCAAGCGGGA	1200
	TGACAGACAG CTAGAGATTT GCAAACAACA GCTCCCAGAT TCTTGCGGCT AGTGGGAGAC	1260
40	CAAGGGACTC CATCGGAGAT NAGGACCCGT TCACAGCTAA GCTRAGCTTT GAGCAAGGTG	1320
40	GCTCARAGTT CGTGCCAGTG GTGGTGAGCC GGCTGGTGCT GCGCTCCATG AGCCGCCGGG	1380
	ATGTCCTCAT CAAGCGATGG CCCCCACCCC TGAGGTGGCA ATACTTCCGC TCACTGCTGC	1440
45	CTGACGCCTC CATTACCATG TGCCCCTCCT GCTTCCAGAT GTTCCATTCT GAGGACTATG	1500
	AGTTGCTGGT GCTTCAGCAT GGCTGCTGCC CCTACTGCCG CAGGTGCAAG GATGACCCTG	1560
50	GCCCATGACC AGCATCCTGG GGACGGCCTG CACCCTCTGC CCGCCTTGGG GTCTGCTGGG	1620
50	CTGTGAAGGA GAATAAAGAG TTAAACTGTC AAAAAAAAAA	1686
	ANAAAAAAA AAAAAAAAA	170

(2) INFORMATION FOR SEQ ID NO: 217:

(i) SEQUENCE CHARACTERISTICS:

-	<ul><li>(A) LENGTH: 999 base pairs</li><li>(B) TYPE: nucleic acid</li><li>(C) STRANDEDNESS: double</li></ul>	
5	(D) TOPOLOGY: linear	
5 ,	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 217:	
	AGCAAATCAC CTTAACGATC TGGAATGAAA CTGTGACCAG TGCCGCCCTG GGTGGTTCTG	60
10	GAGAGACTGC CGTCTTCTTG TTTGGCCATA GGTGCTGGGG CCCCGGCTTC AGTCACTGTC	120
	TCAGACAGKA GTCCCGATAA GCAGATCACC AGTCCTCCAC TGTCCTTCCT GTCGGCCTTG	180
15	CTGCATGAGA AGATAGCTGC TTCCTCCCTC TTTTCCTACA CTGTAAATTA TTGTTTTACA	240
15	ATTGAGTGYC TTAATAATAG TYTACAAATA CTATGTATTT ATGCAAAACT GTTAAAGTTC	300
,	TCATCTGTTA TGATTGGATA CTTGGTCTTG TCAGTAGTGG TCAGCATTGG GTTGTGAGCT	360
20	TGTCCTACTC CATACGTGTT TATCCTGCTA TGCATTTTAC ATTGTGTGTT CACATCTATT	420
	CCAAGGAGCC TTGCTAGAAA CAACACTGGC GGTTCCTGCA GGCCAGGCAG GCATTGGCCC	480
25	ATGCTGTGTC CCATAGGAGC CAATGGAAAG AACGTAGCTT GGTCTGCTAG CCAGCCGTGG	540
23	GGTGGCGCAG GCCAGGCAGC CTCTGCACCA GAGTCCAGCA CCTGCCCATT CCCCAGTCAC	600
	ACAATCATAC TCTTCTTCA TAGAGATTTT ATTACCACCT AGACCACCCT AGTTTTCCTC	660
30	TCTGTTAGTG TCCTGAGCTC TTTTGCAACA AAATGTAGGT ACAGACCAAT CCCTGTCCCT	720
	TCCCCAATCA GGAGCTCCAC ACCATGAGTT GTTTGGTTTT CCAGAAGCTG CCAGTGGGTT	780
35	CCCGTGAATT GCGTTAAGAT ATCGATGATK TTTTTTATTG TTTTTCTTCT TGTTTTTTTA	840
	AATAATATAT TTAAAGGCAG TATCTTTTGT ACTGTGAATT TGCAGTAGAA GATGCAGAAT	900
	GCACTITITI TITACTICIG TIGGIGIGIA TIGIATATAG IGIGIGIGCI TCITGIGAIG	960
40	AAAATAAACT TTTTCTTTAT AAAAAAAAA AAAAAAAA	999
45	(2) INFORMATION FOR SEQ ID NO: 218:	
	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 941 base pairs (B) TYPE: nucleic acid	
50	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 218:	
55	GGCACGAGTA GCATTTCATT TAATCTGCAG GTATATTCTC CCAACAGTTT ATTGTCATGT	60
	GATGTCCTCA GCCAAGATTG TRAGGCAGAG AGGAGCTGTC CCAACCTACT ATACCACCGA	120
	CCCTCCACAG ATCATATTTT TEGTATTAAA CTEGAGTCTC TCCATCCTTC ACATTGTTGA	180

	TGTCCTCTGT	AGCAAACCGG	AAAAGTCAGT	GACAGAAGAT	GCCGCTAGCG	GTTTGAGCCA	240
	GAGAATGACA	GCTCTGGTTT	GGAGAAAAGG	GCCGGATGGT	GGCTCTAGAA	AGCCCATCCT	300
5	TCTGCTCTTC	TTTTTTCTCC	CCCTTATATT	GTGCTTTCAT	TCATTCATTC	ATTCATCAAA	360
	CATTTGTTGA	GCACCTATTA	TGTGTCAAGC	TCTGTGCTAG	CCTCTGGAAA	ACCTGCCCTC	420
10	ATGTAGCTCA	CTGTGGAGTA	GGAGAAACAA	TGACTACACT	ATGATAAGCA	CGGGTTGTCA	480
10	GĠGTCTCACA	GAGCAGTGGC	CCCTCATCCA	GACCGATGAG	GTCAAAGAAG	GCATCCAGGC	540
	GAGGATGGTG	TCAGAGCTAA	CTGAAGAATG	AGAGGGAGCT	GCACCASCAG	GGGTTGGAAC	600
15	TGAAGGTGGC	AGTGCCTGGA	GTCTTGATTC	CAGCAGAGGG	AGAGCAGTCT	GTGAAAAGGC	66
	ACCAAGGGTG	GGAGAGGGCA	GAGCACATGG	AGGAACTTCA	GGTAGTTCTG	GATGGCSCTG	72
00	GGGCAAAGCT	AGAGAGGTAA	GAAGAATCTA	CAAATGTTCC	TCGAGTTACA	TGAACTTCCA	78
20	TCCCAATAAA	CCCATTGGAA	ACGAAAAATT	TAAGTCAĞAA	GIGCATITAA	GGCTGGTCCG	84
	AGTAGAATGA	TTTTTACAAC	GAATTGATCA	CAACCAGTTA	CAGATGTCTT	<b>TGTTCCTTCT</b>	90
25	CCACTCCCAC	TGCTTCACCT	GACTAGCCTI	AAAAAAAT	. A		94

## 30 (2) INFORMATION FOR SEQ ID NO: 219:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 575 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 219:

40	TAAGTGGAAT	CCCCGGGGT	TGCAGGGAAT	TCGGCACGAG	GCATTCTGAG	AAGCTTAAGA	60
	CATACTTIGA	AGACAACCCT	AGGGACCTCC	AGCTGCTGCG	GCATGACCTA	CCTTTGCACC	120
45	CCGCAGTGGT	GAAGCCCCAC	CTGGGCCATG	TTCCTGACTA	CCTGGTTCCT	CCTGCTCTCC	180
45	GTGGCCTGGT	RCGCCCTCAC	AAGAAGCGGA	AGAAGCTGTC	TTCCTCTTGT	AGGAAGGCCA	240
	AGAGAGCAAA	GTCCCAGAAC	CCACTGCGCA	GCTTCAAGCA	CAAAGGAAAG	AAATTCAGAC	300
50	CCACAGCCAA	GCCCTCCTGA	GGTTGTTGGG	CCTCTCTGGA	GCTGAGCACA	TTGTGGAGCA	360
	CAGGCTTACA	CCCTTCGTGG	ACAGGCGAGG	CTCTGGTGCT	TACTGCACAG	CCTGAACAGA	420
e e	CAGTTCTGGG	GCCGGCAGTG	CTGGGCCCTT	TAGCTCCTTG	GCACTTCCAA	GCTGGCATCT	480
55	TGCCCCTTGA	CAACAGAATA	AAAATTTTAG	CTGCCCCAAA		AAAAAAAA	540
	CTCGAGGGGG	GGCCCGTACC	CAATTCGCCC	TATAA		•	575

### (2) INFORMATION FOR SEQ ID NO: 220:

5	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 3018 base pairs
	(B) TYPE: nucleic acid
	(C) STRANDEDNESS: double
	(D) TOPOLOGY: linear

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 220:							•
•	GCCAGCCTTA	CAGGTTTTAC	GTGAAATGAA	AGCCATTGGA	ATAGAACCCT	CGCTTGCAAC	60
15	ATATCACCAT	ATTATTCGCC	TGTTTGATCA	ACCTGGAGAC	CCTTTAAAGA	GATCATCCTT	120
	CATCATTTAT	GATATAATGA	ATGAATTAAT	GGGAAAGAGA	TTTTCTCCAA	AGGACCCGGA	180
20	TGATGATAAG	TTTTTTCAGT	CAGCCATGAG	CATATGCTCA	TCTCTCAGAG	ATCTAGAACT	240
	TGCCTACCAA	GTACATGGCC	TTTTAAAAAC	CGGAGACAAC	TGGAAATTCA	TTGGACCTGA	300
	TCAACATCGT	AATTTCTATT	ATTCCAAGTT	CTTCGATTTG	ATTIGTCTAA	TGGAACAAAT	360
25	TGATGTTACC	TTGAAGTGGT	ATGAGGACCT	GATACCTTCA	GCCTACTTTC	CCCACTCCCA	420
	AACAATGATA	CATCTTCTCC	AAGCATTGGA	TGTGGCCAAT	CGGCTAGAAG	TGATTCCTAA	480
30	AATTTGGGAA	AGATAGTAAA	GAATATGGTC	ATACTTTCCG	CAGTGACCTG	AGAGAAGAGA	540
	TCCTGATGCT	CATGGCAAGG	GACAAGCACC	CACCAGAGCT	TCAGGTGGCA	TTTGCTGACT	600
	GTGCTGCTGA	TATCAAATCT	GCGTATGAAA	GCCAACCCAT	CAGACAGACT	GCTCAGGATT	660
35	GGCCAGCCAC	CTCTCTCAAC	TGTATAGCTA	TCCTCTTTTT	AAGGGCTGGG	AGAACTCAGG	720
	AAGCCTGGAA	AATGTTGGGG	CTTTTCAGGA	AGCATAATAA	GATTCCTAGA	AGTGAGTTGC	780
40	TGAATGAGCT	TATGGACAGT	GCAAAAGTGT	CTAACAGCCC	TTCCCAGGCC	ATTGAAGTAG	840
	TAGAGCTGGC	AAGTGCCTTC	AGCTTACCTA	TTTGTGAGGG	CCTCACCCAG	AGAGTAATGA	900
	GTGATTTTGC	AATCAACCAG	GAACAAAAGG	AAGCCCTAAG	TAATCTAAÇT	GCATTGACCA	960
45	GTGACAGTGA	TACTGACAGC	AGCAGTGACA	GCGACAGTGA	CACCAGTGAA	GGCAAATGAA	1020
	AGTGGAGATT	CAGGAGCAGC	AATGGTCTCA	CCATAGCTGC	TGGAATCACA	CCTGAGAACT	1080
50	GAGATATACC	AATATTTAAC	ATTGTTACAA	AGAAGAAAAG	ATACAGATTT	GGTGAATTTG	1140
	TTACTGTGAG	GTACAGTCAG	TACACAGCTG	ACTTATGTAG	ATTTAAGCTG	CTAATATGCT	1200
	ACTTAACCAT	CTATTAATGC	ACCATTAAAG	GCTTAGCATT	TAAGTAGCAA	CATTGCGGTT	1260
55	TTCAGACACA	TGGTGAGGTC	CATGGCTCTT	GTCATCAGGA	TAAGCCTGCA	CACCTAGAGT	1320
	GTCGGTGAGC	TGACCTCACG	ATGCTGTCCT	CGTGCGATTG	CCCTCTCCTG	CTGCTGGACT	1380
60	TCTGCCTTTG	TTGGCCTGAT	GTGCTGCTGT	GATGCTGGTC	CTTCATCTTA	GGTGTTCATG	1440

	CAGTTCTAAC ACAGTTGGGG TTGGGTCAAT AGTTTCCCAA TTTCAGGATA TTTCGATGTC	1500
	AGAAATAACG CATCTTAGGA ATGACTAAAC AAGATAATGG CAGTTTAGGC TGCACAACTG	1560
5	GTAAAATGAC TGTAGATAAA TGTTGTAATT AGTGTACACG TTTGTATTTT TGTTAATATA	1620
	GCCGCTGCCA TAGTTTTCTA ACTTGAACAG CCATGAATGT TTCATGTCTC CCTTTTTTTT	1680
10	TTGTCTATAG CTGTTACCTA TTTTAGTGGT TGAAATGAGA GCTAGTGATG ACAGAAGGAT	1740
10	GTGGAATGTC TTCTTGACAT CATTGTGTAT TGCTGGTAAT CAAGTTGGTA ACGACTACTT	1800
	CTAGCAGCTC TTACCACTAT GACTTAAGTG GTCCTGGAAG GCAGTAAGTG GAGGTTTGCA	1860
15	GCATTCCTGC CTTCATGAGG GCTTCTACCA CTGACCACTT TGCACGTACC TGGCTCCCAG	1920
	ATTTACTTAG GTACCCCACG AGTCGTCCAC ATAAGCAGCT TCATCTTTAC CTTGCCAGAG	1980
20	TTGACAATTA TGGGATACTC TAGTCTACTT ATACTTGTGT TCCCATCTGT CTGCCATCCT	2040
20	CTGAAGGCCA GGACCCAGTC ATACATCCTT AGAAACCAAA GTATGGTTTT TGTTTTCTCT	2100
	TGGAATGTCA GGTCTTAAGG CATTTAATTG AGGGACAAAA AAAAAAAAA GCCGATATAG	2160
25	TAGCTAGCTA CTTAAGCATC CATGGGTATT GCTCCATATC AAAGCAGATT TGCAGGACAG	2220
	AAAGAGTAAA TTAGCCTTCA GTCTTGGTTT ACAGCTTCCA AGGAGAGCCT TGGSCACCTG	2280
30	AAATGTTAAC TCGGTCCCTT CCTGTCTCTA GTTCATCAGC ACCTGCAGAT GCCTGACTCT	2340
50	TGTTAGCCTT ACTATTCAAT ACAGTCCTTA GATTCACGGT ATGCCTCTTC CTATCCAGGC	2400
	ACCTATTCTG AATCACCATG TIGCTCTGCA GCTAGAGTTG ATAGGAGAAA ATCCATTTGG	2460
35	GTAGATGGCC TATGAATTTG TAGTAGACTT TCAAAATGAG TGATTTGTTA GCTTGGTACT	2520
	TTTAAGTTTG TGGTACAGAT CCTCCAAACC CATACTCTGA GCAATTAACT GCCTTGAACA	2580
40 <sup>-</sup>	TAGAGAAAAA TTAAGGCCTC ACAGGATGAG TCTCCATTCT CTGTAAATGC TTATTTTATC	2640
,,	ATAGTCTTTA GCCTCTAACT ATGAGTAAAA TGTTCTCTTC GGCCGGGTGT GGTGACTCAC	2700
	ACCTGTAACC TCAGCACTTT GGGAGGCAGA GGTGGGAGGA TCACTTAGGT CCAGGAGTTC	2760
45	GAGACTAGCC TGGGCAACAT AGTGAGACAC CGGATCTACA AAAAAATAAA AAGCCAGACT	2820
	GGTGGTATGT ATCTGTGTCC CAGCTAATTG GGAGGGTGAG ATGGGAGGAT TGTTTGAGCC	288
50	TAGGAGAGGG AGGTTGCAGT GAGCCGTGAT CGCACCACTG CACTCCAGCC TGGGCAACAG	294
50	AGCAAGACCC TGTCTTGGAG AAACCAGAAT TTTGGAAGAG CAAATGGGGC TGAGTGCAGT	300
	GGCTCATGCC TGTAATCC	301

<sup>(2)</sup> INFORMATION FOR SEQ ID NO: 221:

. 5	(A) LENGTH: 968 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 221:	
	GGCACGAGGG CCGCGGGACA TCCACGGGGC GCGAGTGACA CGCGGGAGGG AGAGCAGTGT	60
10	TCTGCTGGAG CCGATGCCAA AAACCATGCA TTTCTTATTC AGATTCATTG TTTTCTTTTA	120
	TCTGTGGGGC CTTTTTACTG CTCAGAGACA AAAGAAAGAG GAGAGCACCG AAGAAGTGAA	180
15	AATAGAAGTT TTGCATCGTC CAGAAAACTG CTCTAAGACA AGCAAGAAGG GAGACCTACT	240
13	NAAATGCCCA TTATGACGGC TACCTGGCTA AAGACGGCTC GAAATTCTAC TGCAGCCGGA	300
	CACAAAATGA AGGCCACCCC AAATGGTTTG TTCTTGGTGT TGGGCAAGTC ATAAAAGGCC	360
20	TAGACATTGC TATGACAGAT ATGTGCCCTG GAGAAAAGCG AAAAGTAGTT ATACCCCCTT	420
	CATTTGCATA CGGAAAGGAA GGCTATGCAG AAGGCAAGAT TCCACCGGAT GCTACATTGA	480
25	TTTTTGAGAT TGAACTTTAT GCTGTGACCA AAGGACCACG GAGCATTGAG ACATTTAAAC	540
23	AAATAGACAT GGACAATGAC AGGCAGCTCT CTAAAGCCGA GATAAACCTC TACTTGCAAA	600
	GGGAATTTGA AAAAGATGAG AAGCCACGTG ACAAGTCATA TCAGGATGCA GTTTTAGAAG	660
30	ATATTTTTAA GAAGAATGAC CATGATGGTG ATGGCTTCAT TTCTCCCAAG GAATACAATG	720
	TATACCAACA CGATGAACTA TAGCATATTT GTATTTCTAC TTTTTTTTTT	780
35	CTGTACTTTA TGTATWAAAC AAAGTCMCTT TICTCCMAGT TGKATTTGCT ATTTTTCCCC	840
	TATGAGAAGA TATTTTGATC TCCCCAATAC ATTGATTTTG GTATAATAAA TGTGAGGCTG	900
	TTTTGCAAAC TTAAAAAAAA ATTTAAAAAA ACTGGAGGG GGCCCGTACC CAANTCGCCG	960
40	NATATGAT	968
45	(2) INFORMATION FOR SEQ ID NO: 222:	
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1404 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 222:	
55	CGTTTTCCGG CCGTGCGTTT GTGGCCGTCC GGCCTCCCTG ACATGCAGCC CTCTGGACCC	6
,	CGAGGTTGGA CCCTACTGTG ACACACCTAC CATGCGGACA CTCTTCAACC TCCTCTGGCT	12
60	TGCCCTGGCC TGCAGCCCTG TTCACACTAC CCTGTCAAAG TCAGATGCCA AAAAAGCCGC	18

	CTCAAAGACG	CTGCTGGAGA	AGAGTCAGTT	TTCAGATAAG	CCGGTGCAAG	ACCGGGGTTT	240
	GGTGGTGACG	GACCTCAAAG	CTGAGAGTGT	GGTTCTTGAG	CATCGCAGCT	ACTGCTCGGC	300
5	AAAGGCCCGG	GACAGACACT	TTGCTGGGGA	TGTACTGGGC	TATGTCACTC	CATGGAACAG	360
	CCATGGCTAC	GATGTCACCA	AGGTCTTTGG	GAGCAAGTTC	ACACAGATCT	CACCCGTCTG	420
10	GCTGCAGCTG	AAGAGACGTG	GCCGTGAGAT	GTTTGAGGTC	ACGGGCCTCC	ACGACGTGGA	- 480
10	CCAAGGGTGG	ATGCGAGCTG	TCAGGAAGCA	TGCCAAGGGC	CTGCACATAG	TGCCTCGGCT	540
	CCTGTTTGAG	GACTGGACTT	ACGATGATTT	CCGGAACGTC	TTAGACAGTG	AGGATGAGAT	600
15	AGAGGAGCTG	AGCAAGACCG	TGGTCCAGGT	GGCAAAGAAC	CAGCATTTCG	ATGGCTTCGT	660
	GGTGGAGGTC	TGGAACCAGC	TGCTAAGCCA	GAAGCGCGTG	GCCTCATCC	ACATGCTCAC	720
20	CCACTTGGCC	GAGGCTCTGC	ACCAGGCCCG	GCTGCTGGCC	CTCCTGGTCA	TCCCGCCTGC	780
	CATCACCCC	GGGACCGACC	AGCTGGGCAT	GTTCACGCAC	AAGGAGTTTG	AGCAGCTGGC	840
	CCCCGTGCTG	GATGGTTTCA	GCCTCATGAC	CTACGACTAC	TCTACAGCGC	ATCAGCCTGG	900
25	CCCTAATGCA	CCCCTGTCCT	GGGTTCGAGC	CTGCGTCCAG	GTCCTGGACC	CGAAGTCCAA	960
	GTGGCGAAGC	AAAATCCTCC	TGGGGCTCAA	CTTCTATGGT	ATGGACTACG	CGACCTCCAA	1020
30	GGATGCCCGT	GAGCCTGTTG	TCGGGGCCAG	GTACATCCAG	ACACTGAAGG	ACCACAGGCC	1080
50	CCGGATGGTG	TGGGACAGCC	AGGYCTCAGA	GCACTTCTTC	GAGTACAAGA	AGAGCCGCAG	1140
	TGGGAGGCAC	GTCGTCTTCT	ACCCAACCCT	GAAGTCCCTG	CAGGTGCGGC	TGGAGCTGGC	1200
35	CCGGGAGCTG	GCCTTGGGG	TCTCTATCTG	GGAGCTGGCC	AGGGCCTGGA	CTACTTCTAC	1260
	GACCTGCTCT	AGGTGGGCAT	TGCGGCCTCC	GCGGTGGACG	TGTTCTTTTC	TAAGCCATGG	1320
40	AGTGAGTGAG	CAGGTGTGAA	ATACAGGCCT	NCACTCCGTT	TGCTGTGAAA	АААААААА	1380
.0	АААААААА	АААААААА	AAAA				1404

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### (2) INFORMATION FOR SEQ ID NO: 223:

#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 707 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 223:

55
NGCGCCCTG CAGTCGACAC TAGTGGATCC AAAGAATTCG GCACGAGGGC AGGTCCAGGG 60
CTCAGAAATC AGCTCTATTG ACGAATTCTG CCGCAAGTTC CGCCTGGACT GCCCGCTGGC 120

60 CATGGAGCGG ATCAAGGAGG ACCGGCCCAT CACCATCAAG GACGACAAGG GCAACCTCAA 180

WO 98/54963 PCT/US98/11422

476

-	CCGCTGCATC GCAGACGTGG TCTCGCTCTT CATCACGGTC ATGGACAAGC TGCGCCTGGA	240
5	GATCCGCGCC ATGGATGAGA TCCAGCCCGA CCTGCGAGAG CTGATGGAGA CCATGCACCG	300
3	CATGAGCCAC CTCCCACCCG ACTITGAGGG CCGCCAGACG GTCAGCCAGT GGCTGCAGAC	360
	CCTGAGCGGC ATGTCGGCGT CAGATGAGCT GGACGACTCA CAGGTGCGTC AGATGCTGTT	420
10	CGACCTGGAG TCAGCCTACA ACGCCTTCAA CCGCTTCCTG CATGCCTGAG CCCGGGGCAC	480
	TAGCCCTTGC ACAGAAGGGC AGAGTCTGAG GCGATGGCTC CTGGTCCCCT GTCCGCCACA	540
15	CAGGCCGTGG TCATCCACAC AACTCACTGT CTGCAGCTGC CTGTCTGTG TCTGTCTTTG	600
15	GTGTCAGAAC TTTTGGGCG GGCCCCTCCC CACAATAAAG ATGCTCTCG ACCTTCAAAA	660
	AAAAAAAAA AAAAACTCRG GGGGGGCCCG GTCCCAATCC CCCCNT2;	707
20		
	(2) INFORMATION FOR SEQ ID NO: 224:	
25	(i) SEQUENCE CHARACTERISTICS:  (A) LEXGTH: 1334 base pairs	
	(B) TYPE: nucleic acid	
	(C) STFANDEDNESS: double (D) TOPOLOGY: linear	
30		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 224:	
	GGGGAACTGC AGTGACAGCA GGAGTAAGAG TGGGAGGCAG GACAGAGTTG GGACACAGGT	60
35	ATGGAGAGGG GGTTCAGCGA GCCTAGAGAG GGCAGACTAT CAGGGTGCCG GCGGTGAGAA	120
	TCCAGGGAGA GGAGCGGAAA CAGAAGAGGG GCAGAAGACC GGGGCACTTG TGGGTTGCAG	180
40	AGCCCCTCAG CCATGTTGGG AGCCAAGCCA CACTGGCTAC CAGGTCCCCT ACACAGTCCC	240
	GGGCTGCCCT TGGTTCTGGT GCTTCTGGCC CTGGGGGCCG GGTGGGCTCA GGALGGGTCA	300
	GAGCCCGTCC TGCTGGAGGG GGAGTGCCTG GTGGTCTGTG AGCCTGGCGA AGCTGCTGCA	360
45	GGGGGCCCG GGGGAGCAGC CCTGGGAGAG GCACCCCCTG GGCGAGTGGC ATTTGCTGCG	420
	GTCCGAAGCC AMCACCATGA GCCAGCAGGG GAAACCGGCA ATGGCACCAK TGGGGCCATC	480
50 -	TACTTCGACC AGGTCCTGGT GAACGAGGGC GGTGGCTTTG ACCGGGCTTC TGGCTCCTTC	540
J	GTAGCCCCTG TCCGGGGTGT CTACAGCTTC CGGTTCCATG TGGTGAAGGT GTACAACCGC	600
	CAAACTGTCC AGGTGAGCCT GATGCTGAAC ACGTGGCCTG TCATCTCAGC CTTTGCCAAT	660
55	GATCCTGACG TGACCCGGGA GGCAGCCACC AGCTCTGTGC TACTGCCCTT GGACCCTGGG	720

GACCGAGTGT CTCTGCGCCT GCGTCGGGGG AATCTACTGG GTGGTTGGAA ACACTCAAGT

TTCTCTGGCT TCCTCATCTT CCCTCTCTGA GGACCCAAGT YTTTCAAGCA CAAGAATCCA

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780

WO 98/54963 PCT/US98/11422

477

	GCCCCTGACA	ACTITCTICT	CCCCTCTCTT	GCCCCAGAAA	CAGCAGAGGC	AGGAGAGAGA	900
	CTCCCTCTGG	YTCCTATCCC	ACYTCTTTGC	ATGGGAMCCT	GTGCCAAACA	CCCAAGTTTA	960
5	AGARAARARY.	ARARCTGWGG	CAGGTATACA	GAGCTGGAAG	TGGACCATGG	AAAACATSGA	1020
•	TAACCATGCA	TCYTCTTGCT	TGGCCACCTC	CTGAAACTGT	CCACCTTTGA	AGTTTGAACT	1080
10	TTAGTCCCTC	CAMACTCTGA	CTGCTGCCTC	CTTCCTCCCA	GCTCTCTCAC	TGAGTTATYT	. 1140
	TCACTGTACC	<del>-</del>	TATCCCCACT	ATCTCTCTTT	CTCCTGATCT	GTGCTGTCTT	1200
	ATTCTCCTCC	TTAGGCTTCC	TATTACCTGG	GATTCCATGA	TTCATTCCTT	CAGACCCTCT	1260
15	CCTGCCAGTA	TGCTAAACCC	TCCCTCTCTC	TTTCTTATCC	CGCTGTCCCA	TTGGCCCAGC	1320
	CTGGATGAAT	CTATCAATAA	AACAACTAGA	GAATGGTGGT	САААААААА	AAAAAAAAAC	1380
20	TCGA						1384

(2) INFORMATION FOR SEQ ID NO: 225:

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### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 760 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 225:

GGGTCGACCC ACGCGTCCGC TGACCAGTCC GTTATAGATA CTTCTTCCTA TACCAAAACT 35 GTTTAAACAG GTGCCACCAC AAGGGATGTC GTCCTTACTC TCTGCGGGTC TTCAAGCATC 120 CCTTTGTGGG AAARGTCTCT GGGCAAGCAC GTGGTATTTG GTCTGCTGCT TGCTTCCCTT 180 40 TTTCCACCAG GGATGTTGTG ATCATAAGTC AAAACAACAG TATATTCCAA ATCTCAAAAG 240 CTATTGTGGC CTGAGCACAA TTGAAATCTA GCAGAGTTTT TCCTATGTAG CTTTAGAGTA 300 ACTOTTCTGC TTCTCTGTCA CTTACAATTC AGGITCTGCC TTTGCCTAAG AGCATGAGCA 360 45 GAAGAGTCCT CATGTGACGC TTAGTTCTAT TGCAGTCCTG GGTGAAACTA TTTAAGCWAT GGGGCTGCTK CTCCCCANWT CCTCCCTAAC AATTCGTTGT GTGGACTTCT CATCTAAAAG 480 50 GTTAGTGGCT TTTGCTTGGG ATCAGTGCTC TCTATTGATG TTCTTGCTGG TCTCCAGACA 540 CATTCCTGTT GCATTAAGAC TTGAAAGACT TGTAGATGTG TGATGTTCAG GCACAGGATG 600 CTGAAAGCTA TGTTACTATT CTTAGTTTGT AAATTGTCCT TTTGATACCA TCATCTTGTT 660 55 TTCTTTTGT AGGTATAAAT AAAAACACTG TTGACAATAA AAAAAAAAA AAAAAAAAA 720 ΑΑΑΑΑΑΑΑ ΑΑΑΑΑΑΑΑΑ ΝΑΑΑΑΑΑΑΑ ΑΑΑΑΑΑΑΑΑ 760

(2) INFORMATION FOR SEQ ID NO: 226	226:	NO:	ID	SEQ	FOR	INFORMATION	(2)
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5 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2057 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 226:

	•			<del></del>			
	CCGAGCCGGC	TGCGCCGGGG	GAATCCGTGC	GGGCGCCTTC	CGTCCCRGTC	CCATCCTCGC	60
15	CGCGCTCCAG	CACCTCTGAA	GTTTTGCAGC	GCCCAGAAAG	GAGGCGAGGA	AGGAGGGAGT	120
	GTGTGAGAGG	AGGGAGCAAA	AAGCTCACCC	TAAAACATTT	ATTTCAAGGA	GAAAAGAAAA	180
20	AGGGGGGGCG	CAAAAATGGC	TGGGGCAATT	ATAGAAAACA	TGAGCACCAA	GAAGCTGTGC	240
	ATTGTTGGTG	GGATTCTGCT	CGTGTTCCAA	ATCATCGCCT	TTCTGGTGGG	AGGCTTGATT	300
•	GCTCCAGGGC	CCACAACGGC	AGTGTCCTAC	ATGTCGGTGA	AATGTGTGGA	TGCCCGTAAG	360
25	AACCATCACA	AGACAAAATG	GTTCGTGCCT	TGGGGACCCA	ATCATTGTGA	CAAGATCCGA	420
	GACATTGAAG	AGGCAATTCC	AAGGGAAATT	GAAGCCAATG	ACATCGTGTT	TTCTGTTCAC	480
30 ·	ATTCCCCTCC	CCCACATGGA	GATGAGTCCT	TGGTTCCAAT	TCATGMTGTT	TATCCTGCAG	540
	CTGGACATTG	CCTTCAAGCT	AAACAACCAA	ATCAGRGAAA	ATGCAGAAGT	CTCCATGGAC	600
	GTTTCCCTGG	CTTACCGTGA	TGACGCGTTT	GCTGAGTGGA	CTGAAATGGC	CCATGAAAGA	660
35	GTACCACGGA	AACTCAAATG	CACCTTCACA	TCTCCCAAGA	CTCCAGAGCA	TCGACGCCC	720
	GTTACTATGA	ATGTGATGTC	CTTCCTTTCA	TGGAAATTGG	GTCTGTGGCC	CATGAAGTTT	780
40	TACCTTTTAA	ACATCCGGCT	GCCTGTGAAT	GAGAAGAAGA	AAATCAATGT	GGGAATTGGG	840
	GAGATAAAGG	ATATCCGGTT	GGTGGGGATC	CACCAAAATG	GAGGCTTCAC	CAAGGTGTGG	900
	TTTGCCATGA	AGACCTTCCT	TACGCCCAGC	ATCTTCATCA	TTATGGTGTG	GTATTGGAGG	960
45	AGGATCACCA	TGATGTCCCG	ACCCCCAGTG	CTTCTGGAAA	AAGTCATCTT	TGCCCTTGGG	1020
	ATTTCCATGA	CCTTTATCAA	TATCCCAGTG	GAATGGTTTT	CCATCGGGTT	TGACTGGACC	1080
50	TGGATGCTGC	TGTTTGGTGA	CATCCGACAG	GCATCTTCTA	TGCRATGCTT	CTRTCCTTCT	1140
	GGATCATCTT	CTGTGGCGAG	CACATGATGG	ATCAGCACGA	GCGGAACCAC	ATCGCAGGGT	1200
	ATTGGAAGCA	AGTCGGACCC	ATTGCCGTTG	GTCCTTCTGC	CTCTTCATAT	TTGACATGTG	1260
55	TGAGAGAGGG	GTACAACTCA	CGAATCCCTT	CTACAGTATC	TGGACTACAG	ACATTGGGAA	1320
	CAGAGCTGGC	CATGGCTTTC	ATCATCGTGG	CTGGAATCTG	CCTCTGCCTC	TAACTTCCTG	1380
60	TTTCTATGCT	TCATGGTATT	TCAGGTGTTT	CGGAACATCA	GTGGGAAGCA	GTCCAGCCTG	1440

	CCAGCTATGA	GCAAAGTCCG	GCGGCTACAC	TATGAGGGGC	TAATTTTTAG	GTTCAAGTTC	1500
	CTCATGCTTA	TCACCTTGGC	ansceases	ATGACTGTCA	TOTTOTTCAT	CGTTAGTCAG	1560
5	GTAACGGAAG	GCCATTGGGA	AATGGGGGGG	CGTCACAGTC	CCAAGTGAAC	AGTGCCTTTT	1620
	TCACAGGCAT	CIPLEGGYLE	TGGAATCTGT	YLGICILIEC	TCTGATGTTC	TTGTATGCAC	1680
10	CATCCCATAA	AAACTATGGA	GAAGACCAGT	CCAATGGAAT	GCAACTCCCA	TGTAAATCGA .	1740
10	GGGAAGATTG	recertairs	GTTTCGGAAC	TTTATCAAGA	ATTGTTCAGC	GCTTCGAAAT	1800
	ATTCCTTCAT	CAATGACAAC	ecrecutors	GTATTTGAGT	CAACAAGGCA	ACACATGTTT	1860
15	ATCAGCTTTG	CATTTGCAGT	TOTOLOAGIO	ACATTGATTG	TACTTGTATA	CGCACACAAA	1920
	TACACTGATT	TASCOTTTAT	CTCAAAATGT	TAAATATAAG	GAAAAAGCG	TCAACAATAA	1980
20	ATATICTITG	AGTATTGTCT	TACTTCTCTT	AAAAAAAA	AAAAAACTC	GTGCCGAATT	2040
	CGGCACGAGC	GGEACGA					2057

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### (2) DIFOFMATION FOR SEQ ID NO: 227:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2084 base pairs

(B) TYPE: nucleic acid

(C) STRANCEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 227:

35 GGCAGAGGG CATTTCCTGC AAAGAGCCAA ACCCCCATTC CTCTGTGCCC CTCCTCTCCC 60 ACCARGIGCT TEATARART ASCICITGIT ACCGGRARIA ACTGITCATT TITCACTCCT 120 40 CCCTCCTAGG TCACACTTTT CAGAAAAAA ATCTGCATCC TGGAAACCAG AAGAAAAATA 180 TGAGACGGGG AATCATCGTG TGATGTGTGT SCTGCCTTTG GCTGAGTGTG TGGAGTCCTG 240 CTCAGGTGTT AGGTACAGTG TGTTTGATCG TGGTGGCTTG AGGGGAACCG CTTGTTCAGA 300 45 GOTGTGACTG CGGCTGCACT GCAGAGAAGC TGCCCTTGGC TGCTCGTAGC GCCGGGCCTT 360 CTCTCCTCGT CATCATCCAG ASCAGCCAGT GTCCGGGAGG CAGAAGGTAC CGGGGCAGCT 420 50 ACTGGAGGAD TETGGGGGC TECCTGGGCT GCCCCCTCCG CCGTGGGGCC CTGTTGCTGC 480 TGTCCATCTA TTTCTACTAC TCCCTCCCAA ATGCGGTCGG CCCGCCCTTC ACTTGGATGC 540 TTGCCCTCCT GGGCCTTCTC GCAGGCACTG AACATCCTCC TGGGCCTCAA GGGCCTGGCC 600 55 CCAGCTGAGA TOTOTGCAGT GTGTGAAAAA GGGAATTTCA ACGTGGCCCA TGGGCTGGCA 660 TEGTICATATT ACATOGGATA TOTGCOGOTG ATCOTGCCAG AGCTCCAGGC CCGGATTCGA 720 60 ACTTACAATO AGCATTACAA CAACCTGCTA CGGGGTGCAG TGAGCCAGCG GTGTNATATT 780

	CTCCTCCCAT	TGGACTGTGG	GGTGCCTGAT	AACCTGAGTA	TGGCTGACCC	CAACATTCGC	840
5	TTCCTGGATA	AACTGCCCCA	GCAGACCGGT	GACCGTGCTG	GCATCAAGGA	TCGGGTTTAC	900
5	AGCAACAGCA	TCTATGAGCT	TCTGGAGAAC	GGGCAGCGGG	CGGGCACCTG	TGTCCTGGAG	960
	TACGCCACCC	CCTTGCAGAC	TTTGTTTGCC	ATGTCACAAT	ACAGTCAAGC	TGGCTTTAGC	. 1020
0	GGGGAGGATA	GGCTTGAGCA	GGCCAAACTC	TTCTGCCGGA	CACTTGAGGA	CATCCTGGCA	1080
	GATGCCCCTG	AGTCTCAGAA	CAACTGCCGC	CTCATTGCCT	ACCAGGAACC	TGCAGATGAC	1140
15	AGCAGCTTCT	CGCTGTCCCA	GGAGGTTCTC	CGGCACCTGC	GGCAGGAGGA	AAAGGAAGAG	1200
	GTTACTGTGG	GCAGCTTGAA	GACCTCAGCG	GTGCCCAGTA	CCTCCACGAT	GTCCCAAGAG	1260
	CCTGAGCTCC	TCATCAGTGG	AATGGAAAAG	CCCCTCCCTC	TCCGCACGGA	TTTCTCTTGA	1320
20	GACCCAGGGT	CACCAGGCCA	GAGCCTCCAG	TGGTCTCCAA	GCCTCTGGAC	TGGGGGCTCT	1380
	CTTCAGTGGC	TGAATGTCCA	GCAGAGCTAT	TTCCTTCCAC	AGGGGGCCTT	GCAGGGAAGG	1440
25	GTCCAGGACT	TGACATCTTA	AGATGCGTCT	TGTCCCCTTG	GGCCAGTCAT	TTCCCCTCTC	1500
	TGAGCCTCGG	TGTCTTCAAC	CTGTGAAATG	GGATCATAAT	CACTGCCTTA	CCTCCCTCAC	1560
	CCTTCTTCTC	AGGACTGAGT	GTGTGGAAGT	TTTTCATAAA	CTTTGGATGC	TAGTGTACTT	1620
30	AGGGGGTGTG	CCAGGTGTCT	TTCATGGGGC	CTTCCAGACC	CACTCCCCAC	CCTTCTCCCC	1680
	TTCCTTTGCC	CGGGGACGCC	GAACTCTCTC	AATGGTATCA	ACAGGCTCCT	TCGCCCTCTG	1740
35	GCTCCTGGTC	ATGTTCCATT	ATTGGGGAGC	CCCAGCAGAA	GAATGGAGAG	GAGGAGGAGG	1800
	CTGAGTTTGG	GGTATTGAAT	CCCCGGCTC	CCACCCTGCA	GCATCAAGGT	TGCTATGGAC	1860
	TCTCCTGCCG	GGCAACTCTT	GCGTAATCAT	GACTATCTCT	AGGATTCTGG	CACCACTTCC	1920
40	TTCCCTGGCC	CCTTAAGCCT	AGCTGTGTAT	CGGCACCCCC	ACCCCACTAG	AGTACTCCCT	1980
	CTCACTTGCG	GTTTCCTTAT	ACTCCACCCC	TTTCTÇAACG	GTCCTTTTTT	AAAGCACATC	2040
45	TCAGATTAAA	ААААААААА	ААААААААА	AGGGGGGCN	GCNT		2084

(2) INFORMATION FOR SEQ ID NO: 228:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2143 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 228:

TCGACCCACG CGTCCGGTTG AATTCCTTGA CCTGCAAACA CATATTTATT AGCCTGACTC

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WO 98/54963 PCT/US98/11422

•	AAACAATGAA	GCTATTAAAA	CTTCGGAGGA	ACATTGTAAA	ACTCTCTTTG	TATCGGCATT	120
	TCACCAACAC	GCTTATTTTG	GCAGTGGCAG	CATCCATTGT	GTTTATCATÇ	TGGACAACCA	180
5	TGAAGTTCAG	AATAGTGACA	TGTCAGTCGG	ACTGGCGGGA	GCTGTGGGTA	GACGATGCCA	240
	TCTGGCGCTT	GCTGTTCTCC	ATGATCCTCT	TIGTCATCAT	GGTTCTCTGG	CGACCATCTG	300
10	CAAACAACCA	GAGGTTTGCC	TTTTCACCAT	TGTCTGAGGA	AGAGGAGGAG	GATGAACAAA	. 360
10	AGGAGCCTAT	GCTGAAAGAA	AGCTTTGAAG	GAATGAAAAT	GAGAAGTACC	AAACAAGAAC	420
	CCAATGGAAA	TAGTAAAGTT	AACAAAGCAC	AGGAAGATGA	TTTGAAGTGG	GTAGAAGAGA	480
15	ATGTTCCTTC	TTCTGTGACA	GATGTAGCAC	TTCCAGCCCT	TCTGGATTCA	GATGAGGAAC	540
	GAATGATCAC	ACACTTTGAA	AGGTCCAAAA	TGGAGTAAGG	AATGGGAAGA	TTTGCAGTTA	600
20	AAGATGGCTA	CCATCAGGGA	AGAGATCAGC	ATCTGTGTCA	GTCTTCTGTA	CGGCTCCATG	660
20	GGATTAAAGG	AAGCAATGAC	ATCCTGATCT	GTTCCTTGAT	CTTTGGGCAT	TGGAGTTGGC	720
	GAGAGGTGTC	AGAACAAAGA	GAACATCTTA	CTGAAAACAA	GTTCATAAGA	TGAGAAAAAT	780
25	CTACGAGCTT	CTTATTTACA	ACACTGCTGC	CCCCTTTCCT	CCCAGACTCT	GACATGGATG	840
	TTCATGCAAC	TTAAGTGTGT	TGTTCCTGAA	CTTTCTGTAA	TGTTTCATTT	TTTAAATCTG	900
30	ACAAACTAAA	AAGTTTAACG	TCTTCTAAAA	GATTGTCATC	AACACCATAA	TATGTAATCT	960
,50	CCAGGAGCAA	CTGCCTGTAA	TTTTTATTTA	TTTAGGGAGT	TACATAGGTG	ATGGGGGAAA	1020
	TTGTTAACTA	CCTTTCATTT	TCCTGGGAAG	TCAAGGTTAC	ATCTTGCAGA	GGTTGTTTTG	1080
35	AGAAAAAAGG	GCCCTTCTGA	GTTAAGGAGC	CATAGTTCTA	TCAATGATCA	AAAGAAAAA	1140
	AAAAAAAAGA	GAAACTGTTA	CAGTATGATT	CAGATCATTT	AAAAAAGCAA	AATCAAGTGC	1200
40 -	AATTTTGTTT	ACAAATGGTG	TATATTAAAG	ATTTTTCTAT	TTCAGATGTA	CTTTAAAGAG	1260
	AAATATTAGC	TTAACTCTTT	TGACATCTGC	TATTGTGACA	CATCCCATTG	CTGGCAATGT	1320
	GGTGCACACT	CCGAAACTTT	TAACTACTGT	TTTGTAAGCC	TCCAAGGGTG	GCATTGCAGG	1380
45	GTCCTTAGGC	AATGTTTTGT	TIGCCITTAT	GCAGAGAGGT	GCTCCAAGTG	CTGTGATTGA	144
	GCACCGTGCT	AGAGGAACTG	TAATGCTTCA	GAAGTTGTAG	CTTATACAAA	GGAAACAGGT	150
50	CCTCCTCCCT	TAATTTAAAC	AGTTATTGCA	TGAAGTAGCG	TGGAGGCCCT	GGACTGCTGC	156
50	TCGTTCTTTA	GGATGGACTG	TTCTGGTATC	TGGTATTGGT	TTAGAGACTG	TTAATAAGGG	162
	ACATCACAAG	GTGATGGGAT	TCATTTGAAG	CACTCTATTT	CTGTTTTAAT	GGTTTTATCC	168
55	AATTTTGCCT	TCCCAAGATT	TTTGTTCTAC	ATAAAAAGTT	CATGCCACTT	TTTAATATAA	174
	AAAAATTTAA	CAAAATTAAT	GTATTTTCT	CATTTTTTC	AAACTTTTTC	TAAAGACTCT	180
60	TTCTGTCAAA	CTCATGAAAA	ATTTCTTTCT	ATGGCTTTTA	TTCTAGATTG	TCTTATTTTC	186

WO 98/54963 PCT/US98/11422

482

TGTTAAAACC	AATGACCACA	TGACCACAAT	CTTCACTAAC	TCATACTGCA	GTGAAAGTGT	1920
TAACCCTTAG	GTAGTTTCTC	TACAACTCTT	TGCTATGGTG	ATTTTTAAAA	AAGTTTCCTA	1980
GGGAAGTATC	TCTGAGGGAA	CAGGCAATCT	GAAGGAACTG	ACTATATTCT	CCATGGCTAA	2040
GTCCATTAGG	CCAAAAGNCT	GGGTGGGTAT	TGGTTGTCAN	GCTGTCTATT	GGCATATTAA	2100
AAACGTAGGC	CGGANGGAAT	AATTAGGTTG	TNATGCCGGC	GGG		2143

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#### (2) INFORMATION FOR SEQ ID NO: 229:

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#### (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1025 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

20 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 229:

60 CCTGGCCCAC ATTGCTTCAT TGGCCTGGCC ATGCGCCTGT ACTATGGCAG CCGCTAGTCC 25 CTGACAACTT CCACCCTGAT TCCGGACCCT GTAGATTGGG CGCCACCACC AGATCCCCCT 120 CCCAGGCCTT CCTCCCTCTC CCATCAGCAG CCCTGTAACA AGTGCCTTGT GAGAAAAGCT 180 30 GGAGAAGTGA GGGCAGCCAG GTTATTCTCT GGAGGTTGGT GGATGAAGGG GTACCCTAGG 240 AGATGTGAAG TGTGGGTTTG GTTAAGGAAA TGCTTACCAT CCCCCACCCC CAACCAAGTT 300 CTTCCAGACT AAAGAATTAA GGTAACATCA ATACCTAGGC CTGAGAAATA ACCCCATCCT 360 35 TGTTGGGCAG CTCCCTGCTT TGTCCTGCAT GAACAGAGTT GATGAAAGTG GGGTGTGGGC AACAAGTGGC TTTCCTTGCC TACTTTAGTC ACCCAGCAGA GCCACTGGAG CTGGCTAGTC 480 40 CAGCCCAGCC ATGGTGCATG ACTCTTCCAT AAGGGATCCT CACCCTTCCA CTTTCATGCA 540 600 AGAAGGCCCA GTTGCCACAG ATTATACAAC CATTACCCAA ACCACTCTGA CAGTCTCCTC CAGTTCCAGC AATGCCTAGA GACATGCTCC CTGCCCTCTC CACAGTGCTG CTCCCCACAC 660 45 CTAGCCTTTG TTCTGGAAAC CCCAGAGAGG GCTGGGCTTG ACTCATCTCA GGGAATGTAG 720 CCCCTGGGCC CTGGCTTAAG CCGACACTCC TGACCTCTCT GTTCACCCTG AGGGCTGTCT 780 50 TGAAGCCCGC TACCCACTCT GAGGCTCCTA GGAGGTACCA TGCTTCCCAC TCTGGGGCCT 840 GCCCCTGCCT AGCAGTCTCC CAGCTCCCAA CAGCCTGGGG AAGCTCTGCA CAGAGTGACC 900 TGAGACCAGG TACAGGAAAC CTGTAGCTCA ATCAGTGTCT CTTTAACTGC ATAAGCAATA 960 55 AGATCTTAAT AAAGTCTTCT AGGCTGTAGG GTGGTTCCTA CAACCACAGC CAAAAAAAAA 1020 1025 AAAAA

(2)	INFORMATION	FOR	SEQ	ID	NO:	230:
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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1250 base pairs

(B) TYPE: nucleic acid(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 230:

	GCCCACGCGT	CCGCCCACGC	GTCCGGCGGT	GCGGAGTATG	GGGCGCTGAT	GGCCATGGAG	60
15	GGCTACTGGC	GCTTCCTGGC	GCYGCTGGGG	TCGCCACTGC	TCGTCGGCTT	CCTGTCGGTG	120
	ATSTTCGCCC	TCGTCTGGGT	CCTCCACTAC	CGAGAGGGC	TTGGCTGGGA	TGGGAGCGCA	180
20	CTAGAGTTTA	ACTGGCACCC	AGTGCTSATG	GTCACCGGCT	TCGTCTTCAT	CCAGGGCATC	240
20	GCATCATCGT	CTACAGACTG	CCGTGGACCT	GGAAATGCAG	CAAGCTCCTG	ATGAAATCCA	300
	TCCATGCAGG	GTTAAATGCA	GTTGCTGCCA	TTCTTGCAAT	TATCTCTGTG	CTCCCCCTCT	360
25	TTGAGAACCA	CAATGTTAAC	AATATAGCCA	ATATGTACAG	TCTGCACAGC	TGGGTTGGAC	420
	TGATAGCTGT	CATATGCTAT	TTGTTACAGC	TTCTTTCAGG	TTTTTCAGTC	TTTCTGCTTC	480
30	CATGGGCTCC	GCTTTCTCTC	CGAGCATTTC	TCATGCCCAT	ACATGTTTAT	TCTGGAATTG	540
30	TCATCTTTGG	AACAGTGATT	GCAACAGCAC	TTATGGGATT	GACAGAGAAA	CTGATTTTTT	600
	CCCTGAGAGA	TCCTGCATAC	AGTACATTCC	CGCCAGAAGG	TGTTTTCGTA	AATACGCTTG	660
35	GCCTTCTGAT	CCTGGTGTTC	GGGCCCTCA	TTTTTTGGAT	AGTCACCAGA	CCGCAATGGA	720
	AACGTCCTAA	GGAGCCAAAT	TCTACCATTC	TTCATCCAAA	TGGAGGCACT	GAACAGGGAG	780
40	CAAGAGGTTC	CATGCCAGCC	TACTCTGGCA	ACAACATGGA	CAAATCAGAT	TCAGAGTTAA	840
40	ACARTGAAGT	AGCAGCAAGG	AAAAGAAACT	TAGCTCTGGA	TGAGGCTGGG	CAGAGATCTA	900
	CCATGTAAAA	TGTTGTAGAG	ATAGAGCCAT	ATAACGTCAC	GTTTCAAAAC	TAGCTCTACA	960
45	GTTTTGCTTC	TCCTATTAGC	CATATGATAA	TTGGGCTATG	TAGTATCAAT	ATTTACTTTA	1020
	ATCACAAAGG	ATGGTTTCTT	GAAATAATTT	GTATTGATTG	AGGCCTATGA	ACTGACCTGA	1080
50	ATTGGAAAGG	ATGTGATTAA	TATAAATAAT	AGCAGATATA	AATTGTGGTT	ATGTTACCTT	1140
JU	TATCTTGTTG	AGGACCACAA	CATTAGCACG	GTGCCTTGTG	CAKAATAGAT	ACTCAATATG	1200
	TGAATATGTG	TCTACTAGTA	GTTAATTGGA	TAAACTGGCA	GCATCCCTGA		1250

(2) INFORMATION FOR SEQ ID NO: 231:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1811 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 231:

	CNGNCAGTAC CGGTCNGATT CCCGGGTCGA CCCACGCGTC CGCTGCATTC CAGGGCCTTT	. 60
10	CAGTGGCTTT CATTCTGAAG TTCCTGGATA ACATGTTCCA TGTCTTGATG GCCCAGGTTA	120
	CCASTGTCAT TATCACAACA GTGTCTGTCC TGGTCTTTGA CTTCAGGCCC TCCCTGGAAT	180
1.5	TTTTCTTGGA AGCCSCATCA GTCSTYCTCT CTATATTTAT TTATAATGCC AGCAAGCCTC	240
15	AAGTTCCGGA ATACGCACCT AGGCAAGAAA GGATCCGAGA TCTAAGTGGC AATCTTTGGG	300
	AGCGTTCCAG TGGGGATGGA GAAGAACTAG AAAGACTTAC CAAACCCAAG AGTGATGAGT	360
20	CAGATGAAGA TACTITCTAA CTGGTACCCA CATAGTTTGC AGCTCTCTTG AACCTTATTT	420
	TCACATTITC AGTGTTTGTA ATATTTATCT TTTCACTTTG ATAAACCAGA AATGTTTCTA	480
25	AATCCTAATA TTCTTTGCAT ATATCTAGCT ACTCCCTAAA TGGTTCCATC CAAGGCTTAG	540
25	AGTACCCAAA GGCTAAGAAA TTCTAAAGAA CTGATACAGG AGTAACAATA TGAAGAATTC	600
	ATTAATATCT CAGTACTTGA TAAATCAGAA AGTTATATGT GCAGATTATT TTCCTTGGCC	660
30	TTCAAGCTTC CAAAAAACTT GTAATAATCA TGTTAGCTAT AGCTTGTATA TACACATAGA	720
	GATCAATTTG CCAAATATTC ACAATCATGT AGTTCTAGTT TACATGCCAA AGTCTTCCCT	780
35	TTTTAACATT ATAAAAGCTA GGTTGTCTCT TGAATTTTGA GGCCCTAGAG ATAGTCATTT	840
33	TGCAAGTAAA GAGCAACGGG ACCCTTTCTA AAAACGTTGG TTGAAGGACC TAAATACCTG	900
	GCCATACCAT AGATTTGGGA TGATGTAGTC TGTGCTAAAT ATTTTGCTGA AGAAGCAGTT	960
40	TCTCAGACAC AACATCTCAG AATTITAATT TTTAGAAATT CATGGGAAAT TGGATTTTTG	1020
	TAATAATCTT TTGATGTTTT AAACATTGGT TCCCTAGTCA CCATAGTTAC CACTTGTATT	1080
45	TTAAGTCATT TAAACAAGCC ACGGTGGGGC TTTTTTCTCC TCAGTTTGAG GAGAAAAATC	1140
43	TTGATGTCAT TACTCCTGAA TTATTACATT TTGGAGAATA AGAGGGCATT TTATTTTATT	1200
	AGTTACTAAT TCAAGCTGTG ACTATTGTAT ATCTTTCCAA GAGTTGAAAT GCTGGCTTCA	1260
50	GAATCATACC AGATTGTCAG TGAAGCTGAT GCCTAGGAAC TTTTAAAGGG ATCCTTTCAA	1320
	AAGGATCACT TAGCAAACAC ATGTTGACTT TTAACTGATG TATGAATATT AATACTCTAA	1380
55	AAATAGAAAG ACCAGTAATA TATAAGTCAC TTTACAGTGC TACTTCACAC TTAAAAGTGC	1440
J	ATGGTATTTT TCATGGTATT TTGCATGCAG CCAGTTAACT CTCGTAGATA GAGAAGTCAG	1500
	GTGATAGATG ATATTAAAAA TTAGCAAACA AAAGTGACTT GCTCAGGGTC ATGCAGCTGG	1560
60	GTGATGATAG AAGAGTGGGC TTTAACTGGC AGGCCTGTAT GTTTACAGAC TACCATACTG	1620

PCT/US98/11422

1140

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	TAAATATGAG CTTTATGGTG TCATTCTCAG AAACTTATAC ATTTCTGCTC TCCTTTCTCC	1680
5	TAAGTTTCAT GCAGATGAAT ATAAGGTAAT ATACTATTAT ATAATTCATT TGTGATATCC	1740
3	ACAATAATAT GACTGGCAAG AATTGGTGGA AATTTGTAAT TAAAATAATT ATTAAACCTA	1800
	AAAAAAAN N	. 1811
10		
	(2) INFORMATION FOR SEQ ID NO: 232:	
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 2271 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double	V.
	(D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 232:	
	CTGACCTCAT GGCGTAGAGC CTAGCAACAG CGCAGGCTCC CAGCCGAGTC CGTTATGGCC	60
25	GCTGCCGTCC CGAAGAGGAT GAGGGGCCCA GCACAAGCGA AACTGCTGCC CGGGTCGGCC	120
	ATCCAAGCCC TTGTGGGGTT GGCGCGGCCG CTGGTCTTGG CGCTCCTGCT TGTGTCCGCC	180
30	GCTCTATCCA GTGTTGTATC ACGGACTGAT TCACCGAGCC CAACCGTACT CAACTCACAT	. 240
50	ATTTCTACCC CAAATGTGAA TGCTTTAACA CATGAAAACC AAACCAAACC	300
	CAAATCAGCA CCACCCTCCC TCCCACGACG AGTACCAAGA AAAGTGGAGG AGCATCTGTG	360
35	GTCCCTCATC CCTCGCCTAC TCCTCTGTCT CAAGAGGAAG CTGATAACAA TGAAGATCCT	420
	AGTATAGAGG AGGAGGATCT TCTCATGCTG AACAGTTCTC CATCCACAGC CAAAGACACT	480
40	CTAGACAATG GCGATTATCG AGAACCAGAC TATGACTGGA CCACGGGCCC CAGGGACGAC	540
	GACGAGTCTG ATNGACACCT TGGAAGAAAA CAGGGGTTAC ATGGAAATTG AACAGTCAGT	600
,	GAAATCTTTT AAGATGCCAT CCTCAAATAT AGAAGAGGAA GACAGCCATT TCTTTTTTCA	660
45	TCTTATTATT TTTGCTTTTT GCATTGCTGT TGTTTACATT ACATATCACA ACAAAAGGAA	720
	GATTTTTCTT CTGGTTCAAA GCAGGAAATG GCGTGATGGC CTTTGTTCCA AAACAGTGGA	780
50	ATACCATCGC CTAGATCAGA ATGTTAATGA GGCAATGCCT TCTTTGAAGA TTACCAATGA	840
-	TTATATTTTT TAAAGCACTG TGATTTGAAT TTGCTTATGT AATTTTATTT GCTTGACTTT	900
	TTATATGATA TTGTGCAAAT GTTTGCCATA GGCAATTGGT ACTTAAATGA GAGGTGAGTC	960
55	TCTCTTTTGC CTTGGTGCTT TGGAAATTAA ATGTCACAAA CGAGTATATA ATTTTTTATC	
	TGTACTTTTA GAGCTGAGTT TAATCAGGTG TCCAAAATGT GAGTTAAACA TTACCTTATA	1080

TTTACACTGT TAGTTTTTAT TGTTTTAGAT TTATTATGCT TCTTCTGGAA GTATTAGTGA

PCT/US98/11422

	TGCTACTTTT AAAAGATCCC AAACTTGTAA CTAAATTCTG ACATATCTGT TACTGCTGAC	1200
	TCACATTCAT TCTCCGCCAT TCAAATACTA TTTTTTATCC ACATTTTTT TTGTTCCCAA	1260
5	ACTGTAATGT ACAAGGATAT GTGTGATAAT GCTTTGGATT TGAGTAATAT TTTTTTTCT	1320
	TCCAAGAAAA CTGCTTTGGA TATTTTTAGA TAATTTAAAC ATAATTTAGG ATAATGATAT	1380
10	TGCTCAATCT GACCACAATT TTAGGTAAAA CATTAAATGT GTCAAGAAAT CTTGGCAACA	1440
10	GAGACTCTGC AGCTTGCAGT GGACATAGAT AAAATGTTAC AGAGATACTA TTTTTTTGGT	1500
	TGGAATTACT ATATTAAATT TAGAAGCAGA AACTGGTAAA ATGTTAAATA CATGTACAAT	1560
15	TGCTTTTAGT TAGCAATTGA TTGTAGCATG GGTTCCTCCA AGGTTTCAAG CAATGGGCAG	1620
	AGTTTAAAAT TATATCAGAT TCGTTTACTT CGTTTATTAT TTTACAGTAA ATTTGAATAA	1680
30	ATCTTAGGGG TCATTATCAC TTAAATAATA CTGTACCTAG GTCTTTCAAA TTAAAATTAT	1740
20	ACCTGAATGA AGTTGTTTGT ATACATAAAG GATATTTGTG TACAATTACC TTTTTTCCCC	1800
	CACACTTGTT TTCTTTGTTT TTGTTTTTTA TGGCAACTGG AAAGTATTTA CTATGGGATT	1860
25	CATTTATGTC TGTCTTTCTA TCATAAAGAA TTGATCAATA TGTAAATATG TGATTTGAAC	1920
•	CATGGTTGAC TTACAAGTGT CACTACAGCT TTTTAGAAAA CATAGCCCTA ATATATGTTA	1980
30	AGCAGGACCC GGGTGAGCCA GTGGGCTTGC GCTTTATGTA GAGCTGGAAG AAGGCCGTCC	2040
<i>3</i> 0	ATCCTGTCTC TTGGGCGGAC AGTGTACTTT CCTAATAGGG AAGGGAAGCA CAATGGAAAT	2100
	ACCCCTGAAC CGTTTTATTG CAGTAATTTT TTTCATATCT GAAACTATTA TTTAATATTT	2160
35	TGAATAAGAT TTTAAAAAAT AAATGGCAAA GATATAAATC TAAAAAAAAA AAAAAAAAAA	2220
	N' ANANAAAAA AAAAAAAAA AAAAAAAAA AAAAAAAA	2271
40		
	(2) INFORMATION FOR SEQ ID NO: 233:	
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1338 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 233: 50

	CTTCCGGTTC	TCCGGGCAGC	TGCCACTGCT	GTAGCTTCTG	CCACCTGCCA	CGACCGGGCC	60
55	TCTCCCTGGC	GTTTGGTCAC	CTCTGCTTCA	TTCTCCACCG	CGCCTATGGT	CCCTCTTGGA	120
33	GCCAGCGTGG	CGNGCCTGGC	GGCTCCCGGG	TGGTGAGAGA	CCCCTCCCCC	AACGATGAAG	180
	GCCTCGCAGT	GCTGCTGCTG	TCTCAGCCAC	CTCTTGGCTT	CCGTCCTCCT	CCTGCTGTTG	240
60	CTGCCTGAAC	TAAGCGGGYC	CCTGGMAGTC	CTGCTGCAGG	CAGCCGAGGC	CGCGCCAGGT	300

	YTTGGGCCTC	CTGACCCTAG	ACCAGGACAT	TACCGCCGCT	GCCACCGGGC	CCTWACCCCT	. 360
5	GCCCAGCAGC	CGGCCCTGG	TCTGGCTGAA	GCTGCGGGG	CCGCGGGGCT	CCGAGGGAGG	420
J	CAATGGCAGC	AACCCTGTGG	CCGGGCTTGA	GACGGACGAT	CACGGAGGGA	AGGCCGGGGA	480
	ARGCTCGGTG	GGTGGCGCC	TIGCTGTGAG	CCCCAACCCT	GGCGACAAGC	CCATGACCCA	. 540
10	GCGCGCCCTG	ACCGTGTTGA	TGGTGGTGAG	CGGCGCGGTG	CTGGTGTACT	TCGTGGTCAG	600
	GACGGTCAGG	ATGAGAAGAA	GAAACCGAAA	GACTAGGAGA	TATGGAGTTT	TGGACACTAA	660
15	CATAGAAAAT	ATGGAATTGA	CACCTTTAGA	ACAGGATGAT	GAGGATGATG	ACAACACGTT	720
	GTTTGATGCC	AATCATCCTC	GAAGATAAGA	ATGTGCCTTT	TGATGAAAGA	AÇTTTATCTT	780
	TCTACAATGA	AGAGTGGAAT	TTCTATGTTT	AAGGAATAAG	AAGCCACTAT	ATCAATGTTG	840
20	GGGGGTATT	TAAGTTACAT	ATATTINAAC	AACCTTTAAT	TTGCTGTTGC	AATAAATACC	900
	GTATCCTTTT	ATTATATCTT	TATATGTATA	GAAGTACTCT	GTTAATGGGC	TCAGAGATGT	960
25	TGGGGATAAA	GTATACTGTA	ATAATTTATC	TGTTTGAAAA	ТТАСТАТААА	ACGGTGTTTT	1020
23	CTGRTCGGTT	TTTGTTTCCT	GCTTACCATA	TGATTGTAAA	TIGTTTTATG	TATTAATCAG	1080
	TTAATGCTAA	TTATTTTTGC	TGATGTCATA	TGTTAAAGAG	CTATAAATTC	CAACAACCAA	1140
30	CTGGTGTGTA	AAAATAATTT	AAAATYTCCT	TTACTGAAAG	GTATTTCCCA	TTTTTGTGGG	1200
	GAAAAGAAGC	CAAATTTATT	ACTTTGTGTT	GGGTTTTTA	AAATATTAAG	AAATGTCTAA	1260
35	GTTATTGTTT	GCAAAACAAT	AAATATGATT	TTAAATTCTC	ТТАААААААА	AAAAAAAAAC	1320
<i></i>	CCCGGGGGG	GGCCCGGN					1338

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#### (2) INFORMATION FOR SEQ ID NO: 234:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 31 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 234:

Met Leu Ser Thr Gly Ile Glu Val Ala Arg Pro Pro Ala Thr Leu Leu 50 1 5 10 15

Gly Leu Met Phe Val Leu Thr Gly Met Pro Arg Gly Leu Arg Xaa 20 25 30

55

## (2) INFORMATION FOR SEQ ID NO: 235:

(i) SEQUENCE CHARACTERISTICS:

60 (A) LENGTH: 116 amino acids

					B) TY D) TY	YPE: OPOLO										
			(xi)	SEQU	-					Q II	ои о	: 23	ō:			
5	Met 1	Asn	Val	Val	Ile 5	Val	Ile	Ile	Leu	Phe 10	Ser	Phe	Asp	Ser	Val 15	Gly
10	Thr	Met	Phe	Ser 20	Cys	Asn	Arg	Ile	Pro 25	Lys	Ile	Thr	Val	Leu 30	Asn	Lys
10	Leu	Lys	Phe 35	Xaa	Cys	Glu	Val	Leu 40	Leu	Arg	Ile	Gln	Thr 45	Ile	Gln	Gly
15	Phe	Тут 50	Arg	Cys	Thr	Arg	Ile 55	Ser	Arg	Tyr	Lys	Gly 60	Ile	Phe	Pro	Asp
	Phe 65	Cys	Gln	Ser	Gln	Cys 70	Met	Gly	Cys	Asn	Pro 75	Ģlu	Ser	Xaa	Met	Ala 80
20	Val	Pro	Ala	Leu	Val 85	Thr	Pro	Ile	Leu	Ala 90	His	Arg	Lys	Lys	Glu 95	Lys
25	Gly	Met	Cys	Leu 100	Phe	Thr	Leu	Ile	Ile 105	Ala	Pro	Thr	Arg	Cys 110	Thr	His
	Tyr	Phe	Cys 115	Xaa												
30	(2)	TNE	ODMA	TION	EOD.	SEO	ו מד	vio.	236.							
	(4)	1111		SEQU												
35			(4)	(	A) L B) T		H: 1 ami	.03 a	mino cid		.ds					
			(xi)	SEQ						EQ I	D NO	: 23	6:			
40	Met 1		Ser	Ala	Lys 5		Val	Arg	Gln	Arg 10	Gly	Ala	Val	Pro	Thr 15	Туг
	Tyr	Thr	Thr	Glu 20		Gly	Glu	Ile	Ile 25	Phe	Leu	Val	Leu	Asn 30	Trp	Ser
45	Leu	Ser	Ile 35	Leu	His	Ile	Val	Asp 40		Leu	Суз	Ser	Lys 45		Glu	Lys
50	Ser	Val		Glu	Asp	Ala	Ala 55		Gly	Leu	Ser	Gln 60		Met	Thr	Ala
50	Leu 65		. Trp	Arg	Lys	Gly 70		Asp	Gly	Gly	Ser 75		Lys	Pro	Ile	Leu 80
55	Leu	Lev	ı Phe	Phe	Phe 85		Pro	Leu	Ile	Leu 90		Phe	His	Ser	Phe 95	
•	His	Ser	: Ser	Asn	Ile	Cys	Xaa									

	(2) INFORMATION FOR SEQ ID NO: 237:	
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 42 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 237:</li> </ul>	
10	Met Ile Leu Phe Pro Gln Xaa Ala Leu Arg Leu Gly Xaa Tro Pro Arg 1 5 10 15	
15	Thr Trp Ser Ile Leu Xaa Lys Tyr Ser Val Asn Phe Phe Ser Ala Tyr 20 25 30	
	Ser Pro Met Gly Ala Val Gly Thr Glu Phe 35 40	
20	(2) INFORMATION FOR SEQ ID NO: 238:	
25	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 37 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 238:</li> </ul>	
30	Met Ile Ile Leu Leu Leu Phe Met Leu Leu Asn Asn Val Val Leu Val 1 5 10 15	
	Gln Glu Asp Asn Cys Gln Arg Lys Asn Thr Val Gln Glu Arg Arg Xaa 20 25 30	
35	Trp Ser Gln Trp Xaa 35	
40	(2) INFORMATION FOR SEQ ID NO: 239:	
45	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 128 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 239:</li> </ul>	
50	Met Ala Ala Xaa Pro Pro Gly Cys Thr Pro Pro Xaa Leu Leu Asp Ile 1 5 10 15	<b>&gt;</b>
50	Ser Trp Leu Thr Glu Ser Leu Gly Ala Gly Gln Pro Val Pro Val Glu 20 25 30	1
55	Cys Arg His Arg Leu Glu Val Ala Gly Pro Arg Lys Gly Pro Leu Sex 35 40 45	:
	Pro Ala Trp Met Pro Ala Tyr Ala Cys Gln Arg Pro Thr Pro Leu Thr 50 55 60	:
60	His His Asn Thr Gly Leu Ser Glu Leu Leu Glu His Gly Val Cys Glu	1

	65	•				70					75					80	
~	Glu	Val	Glu	Arg	Val 85	Arg	Arg	Ser	Glu	Arg 90	Tyr	Gln	Thr	Met	Lys 95	Val	
5	Arg	Arg	Ala	Gly 100	Leu	Gly	Pro	Thr	Pro 105	Gly	Met	Ser	Cys	Pro 110	Gly	Asn	
10	Asp	Asn	Thr 115	Val	His	Thr	Met	His 120	Gly	Glu	Ala	Asn	Arg 125	Gly	Ser	Xaa	
15																	
20	(2)	INF	(i)	SEQU ) ) )	ENCE A) L B) T D) T	CHAI ENGT YPE: OPOL E DE:	RACTI H: 6 ami OGY:	ERIS 7 am no a lin	TICS ino cid ear	acid		: 240	):				
25	Met 1	Ser	Ile	Leu	Cys 5	Cys	Pro	Xaa	Leu	Cys 10	Leu	Phe	Phe	Ser	Phe 15	Cys	
30	Ile	Ser	Ser	Gly 20	Ser	Cys	Pro	Phe	Ser 25	His	Val	Ser	Gln	Leu 30	Ser	Phe	
	Ile	Ala	Thr 35		Ser	Gln	Ser	Ser 40		Val	Leu	Leu	Val 45	Pro	Ala	Tyr	
35	Asn	Thr 50		Leu	Ser	Phe	Leu 55	Ala	Phe	Leu	Asp	Cys 60	Ala	Ser	Leu	Thr	
	Ser 65		Xaa														
40	(2)	INF	ORMA	TION	FOR	SEQ	ID 1	NO:	241:								
45	,,,	<del></del>	(i)	SEQU	ENCE (A) I (B) I	CHA LENGI TYPE:	RACT H: 6 ami	ERIS 59 ar .no a	TICS mino acid near	acio		): 24	1:				
50	Met		Thr	Phe	Gln 5		Leu	Leu	. Leu	Ile 10		Ala	Gln	Ser	Thr 15		
55	Lys	: Ile	e Lys	Ser 20		Pro	Leu	His	Met 25		Asn	His	Thr	Leu 30		Asn	
<i>33</i>	Ser	Pro	35 35		a Asr	Pro	Ser	Ser 40		Thr	Leu	Asn	Phe 45		Thr	Gln	
60	Glr	His		ı Ser	· Val	. Ser	Tyr 55		ı Cys	Cys	His	Met 60		Ser	Leu	His	

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His Ala Phe Ala Xaa
      65
5
      (2) INFORMATION FOR SEQ ID NO: 242:
             (i) SEQUENCE CHARACTERISTICS:
10
                    (A) LENGTH: 44 amino acids
                    (B) TYPE: amino acid
                   (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 242:
      Met Val Ser Val Val Leu Ile Phe Ser Phe Leu Ser Leu Thr Ile Ser
15
                       5
      Thr Thr Ala Ser Ala Tyr Asn Gly Asn Asp Thr Gln Gly Trp Asn Asp
20
      Lys Phe His Xaa Xaa Ser Val Lys Thr Gln Thr Xaa
               35
25
      (2) INFORMATION FOR SEQ ID NO: 243:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 51 amino acids
30
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 243:
      Met Ile Ser Asp Ala Gly Ala Gly Phe Gly Val Phe Leu Leu Val Pro
35
      Arg Ala Gly His Cys Trp Gly Ala Gly Lys Pro Leu Pro Ser Cys Pro
      Ser Val Ala Ser Ile Pro Ser Trp Val Leu Pro Ser Phe Leu Glu Arg
40
                                   40
      Gly Arg Xaa
           50
45
       (2) INFORMATION FOR SEQ ID NO: 244:
              (i) SEQUENCE CHARACTERISTICS:
50
                     (A) LENGTH: 43 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 244:
55
       Met Val Gln Thr Ile Gln Asp Phe Leu Ser Leu Phe Ser Thr Pro Ile
                                           10
       Phe Leu Leu Leu Met Phe Glu Thr Leu Ser Leu Ala Pro Ala Trp
 60
                    20
                                       25
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Leu Lys Pro Leu Arg Val Thr Ser His Ser Xaa
                                40
5
     (2) INFORMATION FOR SEQ ID NO: 245:
            (i) SEQUENCE CHAPACTERISTICS:
10
                  (A) LEWFTH: 61 amino acids
                   (3) TYFE: amino acid
                   (D) TOPOLOGY: linear
            (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 245:
15
     Met Ile Leu Met Pro Gly Leu Gly Thr Ser Arg Gln Arg Ser Val Pro
       1 5 10
     Phe Val Pro Thr Leu Ast Ala Ser Thr Pro Gly Ala Met Thr Gly Pro
20
     Thr Ala Thr Leu Thr Ser Cys Glm Tro Thr Thr Ala Cys Arg Val Ser
     Trp Ala Asn Gly Trp Thr Ser Leu Arg Thr Phe Arg Kaa
25
                             55
      (2) INFORMATION FOR SEQ ID NO: 246:
30
             (i) SEQUENCE CHAPACTERISTICS:
                   (A) LEWIH: 36 amino acids
                   (B) TYFE: amino acid
                   (D) TOPOLOGY: linear
35
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 246:
     Met Ser His His Ala Glm Pro Arg Phe Leu Leu Ile Thr Met Leu Leu
             5
                                      10
       1
40
      Gln Glu Ala Lys Pro Val Ser Asn Ile Pro His Leu Leu Glu Ser Trp
                  20
      Tyr Phe Gly Xaa
              35
45
     (2) INFORMATION FOR SEQ ID NO: 247:
50
             (i) SEQUENCE CFARACTERISTICS:
                    (A) LEWIH: 33 amino acids
                    (B) TYFE: amino acid
                    (D) TCFCLOGY: Linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 247:
55
      Met Asn Ser Leu Phe Trp Met Ile Leu Pro Val Ser Gln Asp Gln
                                        10
      Val Val Glu Gly Leu Glm Gly Gly Phe Ser Glm Ile His Met Arg Ile
60
                  20
```

Leu Arg Lys His Leu Xaa 35

5

121	INFORMATION	FOR	SEO	TD	NO:	248:
441	TIME OFFICE TOTAL	1.01	JUQ		110.	4.10.

131	CEOLIEVICE.	CHARACTERTSTICS.

10

- (A) LENGTH: 211 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 248:
- Met Ser Arg Ser Xaa Asp Val Thr Asn Thr Thr Phe Leu Leu Met Ala
  1
  5
  10
  15
  - Ala Ser Ile Tyr Leu His Asp Gln Asn Pro Asp Ala Ala Leu Arg Ala 20 25 30

20

- Leu His Gln Gly Asp Ser Leu Glu Cys Thr Ala Met Thr Val Gln Ile 35 40 45
- Leu Leu Lys Leu Asp Arg Leu Asp Leu Ala Arg Lys Glu Leu Lys Arg 25 50 55 60
  - Met Gln Asp Leu Asp Glu Asp Ala Thr Leu Thr Gln Leu Ala Thr Ala 65 70 75 80
- Trp Val Ser Leu Ala Thr Gly Gly Glu Lys Leu Gln Asp Ala Tyr Tyr 85 90 95
  - Ile Phe Gln Glu Met Ala Asp Lys Cys Ser Pro Thr Leu Leu Leu Leu 100 105 110

35

- Asn Gly Gln Ala Ala Cys His Met Ala Gln Gly Arg Trp Glu Ala Ala 115 120 125
- Glu Gly Leu Leu Gln Glu Ala Leu Asp Lys Asp Ser Gly Tyr Pro Glu 40 130 135 140
  - Thr Leu Val Asn Leu Ile Val Leu Ser Gln His Leu Gly Lys Pro Pro 145 150 155 160
- Glu Val Thr Asn Arg Tyr Leu Ser Gln Leu Lys Asp Ala His Arg Ser 165 170 175
  - His Pro Phe Ile Lys Glu Tyr Gln Ala Lys Glu Asn Asp Phe Asp Arg 180 185 190

- Leu Val Leu Gln Tyr Ala Pro Ser Ala Glu Ala Gly Pro Glu Leu Ser 195 200 205
- Gly Pro Xaa 55 210
  - (2) INFORMATION FOR SEQ ID NO: 249:

			(i) S	( <i>I</i>	A) LI 3) T	ingti (PE :	i: 54 amir	RIST 18 an no ac	nino cid		ds					
5			(xi)	• •				line TION		EQ II	ono:	249	):			
	Met 1	Glu	Asp	Ser	Glu 5	Ala	Leu	Gly	Phe	Glu 10	His	Met	Gly	Leu	Asp 15	Pro
10	Arg	Leu	Leu	Gln 20	Ala	Val	Thr	Asp	Leu 25	Gly	Trp	Ser	Arg	Pro 30	Thr	Leu
15	Ile	Gln	Glu 35	Lys	Ala	Ile	Pro	Leu 40		Leu	Glu	Gly	Lys 45	Asp	Leu	Leu
	Ala	Arg 50	Ala	Arg	Thr	Gly	Ser 55	Gly	Lys	Thr	Ala	Ala 60	Tyr	Ala	Ile	Pro
20	Met 65	Leu	Gln	Leu	Leu	Leu 70	His	Arg	Lys	Ala	Thr 75	Gly	Pro	Val	Val	Glu 80
	Gln	Ala	Val	Arg	Gly 85	Leu	Val	Leu	Val	Pro 90	Thr	Lys	Glu	Leu	Ala 95	Arg
25	Gln	Ala	Gln	Ser 100	Met	Ile	Gln	Gln	Leu 105	Ala	Thr	Tyr	Cys	Ala 110	Arg	Asp
30	Val	Arg	Val 115		Asn	Val	Ser	Ala 120	Ala	Glu	Asp	Ser	Val 125	Ser	Gln	Arg
50	Ala	Val 130		Met	Glu	Lys	Pro 135		Val	Val	Val	Gly 140	Thr	Pro	Ser	Arg
35	Ile 145		Ser	His	Leu	Gln 150		Asp	Ser	Leu	Lys 155		Arg	Asp	Ser	Leu 160
	Glu	Leu	. Leu	Val	Val 165		Glu	. Ala	Asp	Ļeu 170	Leu	Phe	Ser	Phe	Gly 175	Phe
40	Glu	Glu	ı Glu	Leu 180		Ser	Leu	Leu	Cys 185		: Leu	Pro	Arg	Ile 190		Gln
45	Ala	Phe	Leu 195		. Ser	Ala	Thr				ı Asp		Gln 205		Leu	Lys
43	Glu	Le:		. Leu	His	. Asn	215		Thr	Leu	ı Lys	Leu 220		Glu	Ser	Gln
50	Leu 225		Gly	Pro	Asp	Glr 230		ı Glm	Glr	ı Phe	e Glm 235		. Val	. Cys	Glu	Thr 240
	Glu	Gl:	ı Asp	Lys	Phe 245		Leu	ı Lev	тут	250		. Lev	Lys	: Leu	Ser 255	Leu i
55	Ile	e Ar	g Gly	/ Lys 260		. Lev	ı Lev	ı Ph∈	265		n Thr	Lev	ı Glu	270		Tyr
60	Arg	g Le	u Arg 279		ı Phe	Let	ı Glu	ı Glr 280		e Se	r Ile	e Pro	285		s Val	. Leu

WO 98/54963 PCT/US98/11422

	Asn	Gly 290	Glu	Leu	Pro	Leu	Arg 295	Ser	Arg	Cys	His	Ile 300	Ile	Ser	Gln	Phe
5	Asn 305	Gln	Gly	Phe	Tyr	Asp 310	Cys	Val	Ile	Ala	Thr 315	Asp	Ala	Glu	Val	Leu 320
	Gly	Ala	Pro	Val	Lys 325	Gly	Lys	Arg	Arg	Gly 330	Arg	Gly	Pro	Lys	Gly 335	Asp
0	Lys	Ala	Ser	Asp 340	Pro	Glu	Ala	Gly	Val 345	Ala	Arg	Gly	Ile	Asp 350	Phe	His
15	His	Val	Ser 355	Ala	Val	Leu	Asn	Phe 360	Asp	Leu	Pro	Pro	Thr 365	Pro	Glu	Ala
IJ		11e 370	His	Arg	Ala	Gly	Arg 375	Thr	Ala	Arg	Ala	Asn 380	Asn	Pro	Gly	Ile
20	Val 385	Leu	Thr	Phe	Val	Leu 390	Pro	Thr	Glu	Gln	Phe 395	His	·Leu	Gly	Lys	Ile 400
	Glu	Glu	Leu	Leu	Ser 405	Gly	Glu	Asn	Arg	Gly 410	Pro	Ile	Leu	Leu	Pro 415	Tyr
25	Gln	Phe	Arg	Met 420	Glu	Glu	Ile	Glu	Gly 425	Phe	Arg	Tyr	Arg	Cys 430	Arg	Asp
30	Ala	Met	Arg 435	Ser	Val	Thr	Lys	Gln 440	Ala	Ile	Arg	Glu	Ala 445	Arg	Leu	Lys
	Glu	Ile 450		Glu	Glu	Leu	Leu 455		Ser	Glu	Lys	Leu 460		Thr	Tyr	Phe
35	Glu 465		) Asn	Pro	Arg	Asp 470		Gln	Leu	Leu	Arg 475		Asp	Leu	Pro	Leu 480
	His	Pro	Ala	Val	Val 485		Pro	His	Leu	Gly 490		Val	Pro	Asp	Tyr 495	
40	Val	Pro	) Pro	Ala 500		Arg	Gly	Leu	Val 505		Pro	His	Lys	Lys 510		Lys
45	Lys	Leu	515	Ser	Ser	Cys	Arg	520		Lys	Arg		Lys 525		Gln	Asn
-	Pro	530		, Ser	Phe	: Lys	535		Gly	Lys	. Lys	Phe 540		Pro	Thr	Ala
50	Lys 545		Ser	: Xaa	L											
		-												•		
55	(2)	IN	FORM	ATION	FOF	SEÇ	) ID	NO:	250:	•						

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 299 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- 60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 250:

	Met 1	Thr	Thr	Val	Pro 5	Pro	Ser	Pro	Arg	Pro 10	Met	Ser	Arg	Pro	Ser 15	Glu
5	Arg	Asn	Met	Arg 20	Arg	Pro	Arg	Gly	Pro 25	Ser	Pro	Leu	Pro	Ala 30	Ser	Pro
10	Arg	Asn	Ser 35	Thr	Pro	Asp	Glu	Pro 40	Asp	Val	His	Phe	Ser 45	Lys	Lys	Phe
	Leu	Asn 50	Val	Phe	Met	Ser	Gly 55	Arg	Ser	Arg	Ser	Ser 60	Ser	Ala	Glu	Ser
15	Phe 65	Gly	Leu	Phe	Ser	Cys 70	Ile	Ile	Asn	Gly	Glu 75	Glu	Gln	Glu	Gln	Thr 80
	His	Arg	Ala	Ile	Phe 85	Arg	Phe	Val	Pro	Arg 90	His	Glu	Asp	Glu	Leu 95	Glu
20	Leu	Glu	Val	<b>Asp</b> 100	Asp	Pro	Leu	Leu	Val 105	Glu	Leu	Gln	Ala	Glu 110	Asp	Tyr
25	Trp	Tyr	Glu 115	Ala	Tyr	Asn	Met	Arg 120	Thr	Gly	Ala	Arg	Gly 125	Val	Phe	Pro
	Ala	Tyr 130	Tyr	Ala	Ile	Glu <sup>-</sup>	Val 135	Thr	Lys	Glu	Pro	Glu 140	His	Met	Ala	Ala
30	Leu 145	Ala	Lys	Asn	Ser	Asp 150	Trp	Val	Asp	Gln	Phe 155	Arg	Val	Lys	Phe	Leu 160
	Gly	Ser	Val	Gln	Val 165	Pro	Tyr	His	Lys	Gly 170	Asn	Asp	Val	Leu	Cys 175	Ala
35	Ala	Met	Gln	Lys 180	Ile	Ala	Thr	Thr	Arg 185	Arg	Leu	Thr	Val	His 190	Phe	Asn
40	Pro	Pro	Ser 195	Ser	Cys	Val	Leu	Glu 200	Ile	Ser	Val	Arg	Gly 205	Val	Lys	Ile
	Gly	Val 210	Lys	Ala	Asp	Asp	Ser 215	Gln	Glu	Ala	Lys	Gly 220	Asn	Lys	Cys	Ser
45	His 225	Phe	Phe	Gln	Leu	Lys 230	Asn	Ile	Ser	Phe	Cys 235	Gly	Tyr	His	Pro	Lys 240
	Asn	Asn	Lys	Tyr	Phe 245	Gly	Phe	Ile	Thr	Lys 250		Pro	Ala	Asp	His 255	Arg
50	Phe	Ala	Cys	His 260		Phe	Val	Ser	Glu 265		Ser	Thr	Lys	Ala 270	Leu	Ala
55	Glu	Ser	Val 275		Arg	Ala	Phe	Gln 280		Phe	Tyr	Lys	Gln 285	Phe	Val	Glu
55	Tyr	Thr 290	Cys	Pro	Thr	Glu	Asp 295		Tyr	Leu	Glu					

	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: 2	251:							
5			•	(	A) L B) T D) T	ENGT YPE: OPOL	H: 4 ami OGY:	ERIS	ino cid ear	acid		25				
10	Leu 1	Leu	Tyr					PTIO Xaa		_				Ser	Ser 15	Ser
٠	Lys	Gly	Val	Thr 20	Leu	Val	Ser	Met	Asn 25	Leu	Thr	Ser	Phe	Phe 30	Val	Ser
15	Ser	Val	Leu 35	Ala	Cys	Phe	Ser	Xaa 40							•	
20	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: 2	252:							
25			(i) : (xi)	(	A) L B) T D) T	ENGT YPE : OPOL	H: 5 ami OGY:	94 a no a lin	mino cid ear	aci		: 25	2:			
30	Met 1	Pro	Ala	Ser	Ser 5	Leu	Glu	Ser	Arg	Ser 10	Phe	Leu	Leu	Ala	Lys 15	Lys
30	Ser	Gly	Glu	Asn 20	Val	Ala	Lys	Phe	Ile 25	Ile	Asn	Ser	Tyr	Pro 30	Lys	Tyr
35	Phe	Gln	Lys 35	Asp	Ile	Ala	Glu	Pro 40	His	Ile	Pro	Cys	Leu 45	Met	Pro	Glu
	Tyr	Phe 50	Glu	Pro	Gln	Ile	Lys 55	Asp	Ile	Ser	Glu	Ala 60	Ala	Leu	Lys	Glu
40	Arg 65	Ile	Glu	Leu	Arg	Lys 70	Val	Lys	Ala	Ser	Val 75	Asp	Met	Phe	Asp	Gln 80
45	Leu	Leu	Gln	Ala	Gly 85	Thr	Thr	Val	Ser	Leu 90	Glu	Thr	Thr	Asn	Ser 95	Leu
			Xaa	100					105					110	_	_
50			Gln 115					120					125			
<i></i>		130	Ser				135					140				
55	145		Asn			150					155					160
60	Glu	His	Ser	Tyr	Cys 165	Thr	Met	Ile	Arg	Gly 170	Met	Val	Lys	His	Arg 175	Ala

	Tyr	GIu	Gin	180	Leu	Asn	Leu	ıyr	185	GIU	Leu	Leu	ASN	190	Arg	Leu
5	His	Ala	Asp 195	Val	Tyr	Thr	Phe	Asn 200	Ala	Leu	Ile	Glu	Ala 205	Thr	Val	Cys
	Ala	Ile 210	Asn	Glu	Lys	Phe	Glu 215	Glu	Lys	Trp	Ser	Lys 220	Ile	Leu	Glu	Leu
10	Leu 225	Arg	His	Met	Val	Ala 230	Gln	Lys	Val	Lys	Pro 235	Asn	Leu	Gln	Thr	Phe 240
15	Asn	Thr	Ile	Leu	Lys 245	Cys	Leu	Arg	Arg	Phe 250	His	Val	Phe	Ala	Arg 255	Ser
	Pro	Ala	Leu	Gln 260	Val	Leu	Arg	Glu	Met 265	Lys	Ala	Ile	Gly	Ile 270	Glu	Pro
20	Ser	Leu	Ala 275	Thr	Tyr	Ĥis	His	Ile 280	Ile	Arg	Leu	Phe	Asp 285	Gln	Pro	Gly
	Asp	Pro 290	Leu	Lys	Ārg	Ser	Ser 295	Phe	Ile	Ile	Tyr	Asp 300	Ile	Met	Asn	Glu
25	<b>L</b> eu 305	Met	Gly	Lys	Arg	Phe 310	Ser	Pro	Lys	Asp	Pro 315	Asp	Asp	Asp	Lys	Phe 320
30	Phe	Gln	Ser	Ala	Met 325	Ser	Ile	Cys	Ser	Ser 330	Leu	Arg	Asp	Leu	Glu 335	Leu
	Ala	Tyr	Gln	Val 340	His	Gly	`Leu	Leu	Lys 345	Thr	Gly	Asp	Asn	Trp 350	Lys	Phe
35	Ile	Gly	Pro 355	Asp	Gln	His	Arg	Asn 360	Phe	Tyr	Tyr	Ser	Lys 365	Phe	Phe	Asp
	Leu	Ile 370		Leu	Met	Glu	Gln 375	Ile	Asp	Val	Thr	Leu 380	Lys	Trp	Tyr	Glu
40	Asp 385	Leu	Ile	Pro	Ser	Ala 390	Tyr	Phe	Pro	His	Ser 395	Gln	Thr	Met	Ile	His 400
45	Leu	Leu	Gln			Asp				Arg 410		Glu	Val	Ile	Pro 415	Lys
	Ile	Trp	Lys	Asp 420		Lys	Glu	Tyr	Gly 425		Thr	Phe	Arg	Ser 430		Leu
50 ·	Arg	Glu	Glu 435		Leu	Met	Leu	Met 440		Arg	Asp	Lys	His 445		Pro	Glu
	Leu	Gln 450		Ala	Phe	Ala	Asp 455		Ala	Ala	Asp	Ile 460		Ser	Ala	Tyr
55	Glu 465		Gln	Pro	Ile	Arg 470		Thr	Ala	Gln	Asp 475		Pro	Ala	Thr	Ser 480
60	Leu	Asn	Cys	Ile	Ala 485		Leu	Phe	Leu	Arg 490		. Gly	Arg	Thr	Gln 495	Glu.

•	Ala	Trp	Lys	Met 500	Leu	Gly	Leu	Phe	Arg 505	Lys	His	Asn	Lys	Ile 510	Pro	Arg
5	Ser	Glu	Leu 515	Leu	Asn	Glu	Leu	Met 520	Asp	Ser	Ala	Lys	Val 525	Ser	Asn	Ser
	Pro	Ser 530	Gln	Ala	Ile	Glu	Val 535	Val	Glu	Leu	Ala	Ser 540	Ala	Phe	Ser	Leu
10	Pro 545	Ile	Cys	Glu	Gly	Leu 550	Thr	Gln	Arg	Val	Met 555	Ser	Asp	Phe	Ala	Ile 560
15	Asn	Gln	Gļu	Gln	Lys 565	Glu	Ala	Leu	Ser	Asn 570	Leu	Thr	Ala	Leu	Thr 575	Ser
	Asp	Ser	Asp	Thr 580	Asp	Ser	Ser	Ser	Asp 585	Ser	Asp	Ser	Asp	Thr 590	Ser	Glu
20	Gly	Lys												,		
25	(2)	INF		TION SEQU	ENCE	СНА	RACT	ERIS	TICS		đe					
					-				mino	acı	as					
30			(xi)	•	D) 1	YPE: OPOL E DE	OGY:	lin	ear	EQ I	D NO	: 25	3 :			
30	Met 1	Lys		(	D) I	OPOL E DE Cys	OGY: SCRI	lin PTIO	ear N: S					Pro	Leu 15	Leu
30 35	1		Leu	SEQ	D) I UENC Leu 5	OPOL E DE Cys	OGY: SCRI	lin PTIO Pro	ear N: S Asn	Trp 10	Ala	Arg	Cys		15	
35	1 Leu	Leu	Leu Phe	SEQ Asn Pro 20 Lys	D) T UENC Leu 5 Gln	OPOL E DE Cys Leu	OGY: SCRI Ile Leu	lin PTIO Pro	ear N: S Asn Phe 25	Trp 10 Gln	Ala	Arg Glu	Cys	Asp 30	15 Asp	Pro
	1 Leu Leu	Leu Lys	Phe Ala 35	SEQ Asn Pro 20 Lys	D) TUENC Leu 5 Gln	OPOL E DE Cys Leu Ala	OGY: SCRI Ile Leu Asn	Pro Pro Leu 40	N: S Asn Phe 25 Val	Trp 10 Gln Glu	Ala Gly Ala	Arg Glu Val	Cys Asp Pro 45	Asp 30 Trp	15 Asp Gly	Pro
35	1 Leu Leu	Leu Lys Ala 50	Phe Ala 35	SEQ Asn Pro 20 Lys	D) TUENC  Leu  5  Gln  Ala	Cys Leu Ala	OGY: SCRI Ile Leu Asn Val	lin PTIO Pro Pro Leu 40	N: S Asn Phe 25 Val	Trp 10 Gln Glu Leu	Ala Gly Ala Val	Arg Glu Val Arg 60 Ala	Cys Asp Pro 45 Val	Asp 30 Trp Gln	15 Asp Gly Leu	Pro Ile Gln
35	Leu Leu Lys Ser 65	Leu Lys Ala 50 Cys	Phe Ala 35 Pro	SEQ Asn Pro 20 Lys	D) I UENC Leu 5 Gln Ala Phe	OPOLICE DE CYS  Leu  Ala  Gln  Arg  70	OGY: SCRI Ile Leu Asn Val 55	Pro Pro Leu 40 Thr	ear N: S Asn Phe 25 Val Cys	Trp 10 Gln Glu Leu	Ala Gly Ala Val Leu 75 Ser	Arg Glu Val Arg 60 Ala	Cys Asp Pro 45 Val	Asp 30 Trp Gln Ser	Asp Gly Leu Gln	Pro Ile Gln Ser 80 Val
35	Leu Lys Ser 65	Leu Lys Ala 50 Cys	Phe Alaa 35 Pro	( SEQ Asn Pro 20 Lys Ser	D) I UENC Leu 5 Gln Ala Phe Ser 85	OPOL E DE Cys Leu Ala Gln Arg 70	OGY: SCRI Ile Leu Asn Val 55	lim PTIO Pro Pro Leu 40 Thr	N: S Asn Phee 25 Val Cys Thr	Trp 10 Glu Leu Leu Met	Ala Gly Ala Val Leu 75	Arg Glu Val Arg 60 Ala	Cys Asp Pro 45 Val Thr	Asp 30 Trp Gln Ser	Asp Gly Leu Gln Pro 95	Pro Ile Gln Ser 80 Val
35 40 45	Leu Lys Ser 65 Pro	Leu Lys Alaa 50 Cys	Phe Ala 35 Pro	Asn Pro 20 Lys Ser Pro Ile 100	D) I UENC Leu 5 Gln Ala Phe Ser 85	COPOL E DE Cys Leu Ala Gln 70 Cys	OGY: SCRI Ile Leu Asn Val 55 Pro	lim PTIO Pro Pro Leu 40 Thr	Phe 25 Val Cys Thr Pro Val 105	Trp 10 Glu Leu Leu Met	Ala Gly Ala Val Leu 75 Ser	Arg Glu Val Arg 60 Ala His	Cys Asp Pro 45 Val Thr	Asp 30 Trp Gln Ser Pro Gln 110	Asp Gly Leu Gln Pro 95	Pro Ile Gln Ser 80 Val

-	(2)	INFO	RMAT	ION	FOR	SEQ	ID N	10: 2	54:							
5		٠	(i) S (xi)	() (I)	A) LI 3) T? O) T(	ENGTI (PE: OPOLA	f: 2: amir XGY:	l am no ac line	ino a cid ear	acid		: 254	<b>1</b> :			
10	Met 1	Arg	Tyr	His	Ala 5	Gln	Leu '	Ile	Phe	Cys 10	Ile	Phe	Суз	Xaa	Phe 15	Val
	Phe	Val	Xaa	Lys 20	Xaa											
15																
	(2)	INFO	ORMAT	NOI	FOR	SEQ	ID N	10: 2	255:							
20			(i) 5 (xi)	() () ()	A) LI B) T D) T	ENGT YPE: OPOL	H: 3: ami: OGY:	l am no a lin	ino a cid ear	acid		: 255	5:			
25	Met 1	Asn	Asp	Asn	Ser 5	Pro	Asn	His	Ser	Ser 10	Ser	Tyr	Leu	Pro	Leu 15	Pro
30	Leu	Thr	Ile	Val 20	Ile	Leu	Gln	Thr	Gly 25	His ,	Lys	Gly	Thr	Leu 30	Xaa	•
	(2)	INF	ORMAT	CION	FOR	SEQ	ID 1	VO: 2	256:							
35			(i) :	· (:	A) L B) T	engt Ype:	H: 2	ERIS 19 a no a lin	mino cid		ds					
40			(xi)							EQ II	ON C	: 25	б:			
	Met 1	His	Phe	Leu	Phe 5	Arg	Phe	Ile	Val	Phe 10	Phe	Tyr	Leu	Trp	Gly 15	Leu
45	Phe	Thr	Ala	Gln 20	Arg	Gln	Lys	Lys	Glu 25	Glu	Ser	Thr	Glu	Glu 30	Val	Lys
	Ile	Glu	Val 35	Leu	His	Arg	Pro	Glu 40	Asn	Суз	Ser	Lys	Thr 45	Ser	Lys	Lys
50	Gly	Asp 50	Leu	Leu	Asn	Ala	His 55	Tyr	Asp	Gly	Tyr	Leu 60	Ala	Lys	Asp	Gly
55	Ser 65	Lys	Phe	Туг	Cys	Ser 70	Arg	Thr	Gln	Asn	Glu 75	Gly	His	Pro	Lys	Trp 80
<i>33</i>	Phe	Val	Leu	Gly	Val 85	Gly	Gln	Val	Ile	Lys 90	Gly	Leu	Asp	Ile	Ala 95	Met
60	Thr	Asp	Met	Cys 100	Pro	Gly	Glu	Lys	Arg 105	Lys	Val	Val	Ile	Pro 110	Pro	Ser

	Phe	Ala	Tyr 115	Gly	Lys	Glu	Gly	Tyr 120	Ala	Glu	Gly	Lys	11e 125	Pro	Pro	Asp
5	Ala	Thr 130	Leu	Ile	Phe	Glu	Ile 135	Glu	Leu	Tyr	Ala	Val 140	Thr	Lys	Gly	Pro
10	Arg 145	Ser	Ile	Glu	Thr	Phe 150	Lys	Gln	Ile	Asp	Met 155	Asp	Asn	Asp	Arg	Gln 160
	Leu	Ser	Lys	Ala	Glu 165	Ile	Asn	Leu	Tyr	Leu 170	Gln	Arg	Glu	Phe	Glu 175	Lys
15	Asp	Glu	Lys	Pro 180	Arg	Asp	Lys	Ser	Туг 185	Gln	Asp	Ala	Val	Leu 190	Glu	Asp
	Ile	Phe	Lys 195	Lys	Asn	Asp	His	Asp 200	Gly	Asp	Gly	Phe	Ile 205	Ser	Pro	Lys
20	Glu	Tyr 210	Asn	Val	Tyr	Gln	His 215	Asp	Glu	Leu	Xaa					
25	(2)	INFO	ORMAT	MOTO	FOR	SEO	י מד	io :	57.			,				
	(-,		(i) :							:						
30			(xi)	C	B) T D) T	YPE: OPOL	ami OGY:	no a	cid ear	acid		. 25°	7		-	
	Met		Val											Phe	Val	Leu
35	1				5					10					15	•
	Phe	Trp	Ser	Val 20	His	Cys	Ile	Ser	Asp 25	Lys	Phe	Gly	Cys	Leu 30	Trp	His
40	Val	Cys	Met 35	Lys	Arg	Glu	Gly	Asp 40	Xaa	Asn	Cys		Ser 45	Phe	Ser	Xaa
-	Leu	<b>Xaa</b> 50									·	•				
45															. •	
	(2)	INF	ORMAI	CION	FOR	SEQ	ID N	IO: 2	:58							
50			(i)	() ()	A) Li B) T D) T	ENGT YPE: OPOL	H: 1: ami: OGY:	22 an no ao line	mino cid ear	aci		: 258	3:			
55	Met 1	Pro	Ser	Gln	Thr 5	Glu	Xaa	Phe	Ala	Ala 10	Cys	Gly	Gly	His	Ser 15	Leu
60	Leu	Leu	Val	Xaa 20	Leu	Pro	Leu	Gly	Leu 25	Pro	Phe	Cys	Pro	Arg 30	Ala	Ala.

	Leu	Cys	Asp 35	Leu	Pro	Phe	Ser	Leu 40	Pro	Ser	Phe	Pro	Gly 45	Gln	Ala	Arg
5	Arg	Gly 50	Gly	Ala	Glu	Lys	Gln 55	Gly	Ala	Glu	Gly	Arg 60	Gly	Leu	Gln	Val
	Lys 65	Pro	Arg	Gly	Gln	Arg 70	Thr	Phe	Gln	Val	Ser 75	Arg	Thr	Ala	Pro	Ala 80
10	Ala	Pro	Arg	Ser	Arg 85	Gln	Pro	Arg	Pro	Pro 90	Ala	Ala	Leu	Pro	Ala 95	Leu
15	Gly	Phe	Gly	Gly 100	Arg	Gly	Val	Ala	Lys 105	Gly	Arg	Phe	Leu	Cys 110	Phe	Trp
	Cys	Leu	Туг 115	Met	Leu	Arg	Ile	Asp 120	Gln	Xaa						
20	(2)	INF	ORMA'	TION	FOR	SEQ	ID	NO: I	259:						:	
25			(i) (xi)	(	A) I B) I D) I	ENGT YPE : OPOL	H: 8 ami OGY:	ERIS 8 am no a lin PTIO	ino cid ear	acid		; 25	9:			
30	Met		Ala	Phe	Cys 5		Leu	Leu	Leu	Gln 10	Ala	Gln	Ser	Leu	Leu 15	Pro
	Arg	Thr	Met	Ala 20		Pro	Gln	Asp	Ser 25		Arg	Prò	Gly	Glu 30	Glu	Asp
35	Glu	Gly	Met 35		Leu	Leu	Gln	Thr 40		Asp	Ser	Met	Ala 45		Gly	Ala
40	Arg	Pro 50		Ala	Xaa	Arg	Gly 55		Ala	Arg	Trp	Gly 60		Ala	Tyr	Thr
	Leu 65		His	Asn	Pro	Thr 70		Gln	(Val	. Phe	Arg 75		Thr	Ala	Leu	80
45	Gly	Ala	Asn	Gly	Ala 85		Pro	Хаа	<b>L</b>							
		.•														
50	(2)	IN	FORMA			_	-	NO: Teris								
•			•		(A)	LENG	TH:	26 au ino a	mino		ds					
55			(xi		(D)	ropo:	LOGY	: li:	near	SEQ :	ID NO	): 20	50:			
		z Ile L	e Glr	ı Val	l Sex		l Pro	) Le	ı Lev	ı Thr 10		e Met	: Ile	e Phe	Leu 19	Leu
60	Тут	r Le	u Gli	n Ile	e Gly	/ Pro	Gly	, Lys	. Let	ı Xaa	ı					

WO 98/54963

503

5	(2) INFORMATION FOR SEQ ID NO: 261:
10	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 29 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 261:</li> </ul>
15	Met Leu Leu Asp Pro Phe Ile Leu Leu Phe Cys Leu Phe Ser Thr Ala 1 5 10 15  Ala Gln Ser Cys Leu Glu Phe Ile Tyr Ile Gln Phe Xaa 20 25
20	(2) INFORMATION FOR SEQ ID NO: 262:
25 .	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 44 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 262:
30	Met Lys Phe Leu Ser Ile Leu Leu Asp Asp Asn Asn Phe Xaa Leu Met  1 5 10 15
	Leu Met Leu Ala Pro Phe Gly Cys Leu Ala Phe Glu Arg Ser Met Lys 20 25 30
35	Met Arg Asn Gly Ala Leu Gly Leu Glu Glu Val Xaa 35 40
40	(2) INFORMATION FOR SEQ ID NO: 263:
45	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 363 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 263:</li> </ul>
50	Met Arg Thr Leu Phe Asn Leu Leu Trp Leu Ala Leu Ala Cys Ser Pro 1 5 10 15
	Val His Thr Thr Leu Ser Lys Ser Asp Ala Lys Lys Ala Ala Ser Lys 20 . 25 . 30
55	Thr Leu Leu Glu Lys Ser Gln Phe Ser Asp Lys Pro Val Gln Asp Arg 35 40 45
	Gly Leu Val Val Thr Asp Leu Lys Ala Glu Ser Val Val Leu Glu His 50 55 60
60	Arg Ser Tyr Cys Ser Ala Lys Ala Arg Asp Arg His Phe Ala Gly Asp

	65					70					75					80
5	Val	Leu	Gly	Тут	Val 85	Thr	Pro	Trp	Asn	Ser 90	His	Gly	Tyr	Asp	Val 95	Thr
J	Lys	Val	Phe	Gly 100	Ser	Lys	Phe	Thr	Gln 105	Ile	Ser	Pro	Val	Trp 110	Leu	Gln
10	Leu	Lys	Arg 115	Arg	Gly	Arg	Glu	Met 120	Phe -	Glu	Val	Thr	Gly 125	Leu	His	Asp
	Val	Asp 130	Gln	Gly	Trp	Met	Arg 135	Ala	Val	Arg	Lys	His 140	Ala	Lys	Gly	Leu
15	His 145	Ile	Val	Pro	Arg	Leu 150	Leu	Phe	Glu	Asp	Trp 155	Thr	Tyr	Asp	Asp	Phe 160
20	Arg	Asn	Val	Leu	Asp 165	Ser	Glu	Asp	Glu	Ile 170	Glu	Glu	Leu	Ser	Lys 175	Thr
	Val	Val	Gln	Val 180	Ala	Lys	Asn	Gln	His 185	Phe	Asp	Gly	Phe	Val 190	Val	Glu
25	Val	Trp	Asn 195	Gln	Leu	Leu	Ser	Gln 200	Lys	Arg	Val	Thr	Asp 205	Gln	Leu	Gly
	Met	Phe 210	Thr	His	Lys	Glu	Phe 215	Glu	Gln	Leu	Ala	Pro 220	Val	Leu	Asp	Gly
30	Phe 225	Ser	Leu	Met	Thr	Туг 230	Asp	Tyr	Ser	Thr	Ala 235	His	Gln	Pro	Gly	Pro 240
35	Asn	Ala	Pro	Leu	Ser 245	Trp	Val	Arg	Ala	Суs 250	Val	Gln	Val	Leu	Asp 255	Pro
	Lys	Ser	Lys	Trp 260	Arg	Ser	Lys ·	Ile	Leu 265	Leu	Gly	Leu	Asn	Phe 270	Туг	Gly
40	Met	Asp	Тут 275	Ala	Thr	Ser	Lys	Asp 280	Ala	Arg	Glu	Pro	Val 285	Val	Gly	Ala
	Arg	Tyr 290	Ile	Gln	Thr	Leu	Lys 295	Asp	His	Arg	Pro	Arg 300	Met	Val	Trp	Asp
45	Ser 305	Gln	Xaa	Ser	Glu	His 310	Phe	Phe	Glu	Tyr	Lys 315	Lys	Ser	Arg	Ser	Gly 320
50	Arg	His	Val	Val	Phe 325	Tyr	Pro	Thr	Leu	Lys 330	Ser	Leu	Gln	Val	Arg 335	Leu
	Glu	Leu	Ala	Arg 340	Glu	Leu	Gly	Val	Gly 345	Val	Ser	Ile	Trp	Glu 350	Leu	Gly
<b>5</b> 5	Gln	Gly	Leu 355	Asp	Tyr	Phe	Tyr	Asp 360	Leu	Leu	Xaa					

WO 98/54963 PCT/US98/11422

			(i)	(	A) I B) 1	CHA ENGI YPE:	H: 1	.28 a	mino cid		.ds					
5			(xi)	SEQ						EQ I	D NO	: 26	4.:			
	Leu 1		Thr	Lys	Ile 5	Leu	Val	Lys	Pro	Asp 10	Arg	Thr	Phe	Glu	Ile 15	Lys
10	Ile	Gly	Gln	Pro 20	Thr	Val	Ser	Tyr	Phe 25	Leu	Lys	Ala	Ala	Ala 30		Ile
15	Glu	Lys	Gly 35	Ala	Arg	Gln	Thr	Gly 40	Lys	Glu	Val	Ala	Gly 45	Leu	Val	Thr
	Leu	Lys 50	His	Val	Tyr	Glu	Ile 55	Ala	Arg	Ile	Lys	Ala 60	Gln	Asp	Glu	Ala
20	Phe 65		Leu	Gln	Asp	Val 70	Pro	Leu	Ser	Ser	Val 75	Val	Arg	Ser	Ile	Ile 80
	Gly	Ser	Ala	Arg	Ser 85	Leu	Gly	Ile	Arg	Val 90	Val	Lys	Asp	Leu	Ser 95	Ser
25	Glu	Glu	Leu	Ala 100	Ala	Phe	Gln	Lys	Glu 105	Arg	Ala	Ile	Phe	Leu 110	Ala	Ala
30	Gln	Lys	Glu 115	Ala	Asp	Leu	Ala	Ala 120	Gln	Glu	Glu	Ala	Ala 125	Lys	Lys	Xaa
35	(2)	· INF	ORMAT	rion	FOR	SEQ	ID 1	NO: 2	265 :							
40			(i) :	(1 (1	A) L B) T D) T	CHAI ENGT YPE: OPOL E DE:	H: 5 ami: OGY:	4 am no a lin	ino d cid ear	acid		: 26!	5:			
45	Met 1	Leu	Leu	Gln	Ile 5	His	Pro	Leu	Leu	Pro 10	Ser	Pro	Thr	Ile	Pro 15	His
	Ile	Leu	Leu	Leu 20	Phe	Leu	Tyr	Pro	Thr 25	Phe	Ser	Ile	Leu	Glu 30	His	Ser
50	Cys	Ser	Tyr 35	Cys	Ile	Glu	Tyr	Leu 40	Trp	Val	Cys	Leu	Leu 45	Phe	Cys	Leu
55	Ser	Leu 50	Trp	Phe	Leu	Xaa										

(2) INFORMATION FOR SEQ ID NO: 266:

60 (i) SEQUENCE CHARACTERISTICS:

				(	(A) L	ENG	'H: 2	:9 an	uno	acio	IS					
				(	(B) I	YPE:	ami	.no a	cid							
								lir								
_			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 26	6:			
5											•					
	Met	Суѕ	Leu	Trp	Cys	Cys	Gly	Asp	Val	Cys	Ser	Gly	Leu	Ser	Ser	Let
	1				5					10					15	
																•
	Leu	Ser	Leu	Cys	Val	Cys	Cys	Val	Val	Leu	Ala	Val	Cys			
10				20					25				_			
•																
	(2)	INF	ORMA'	TION	FOR	SEO	ID I	NO:	267:							
15						_										
			(i)	SEOU	ENCE	СНА	RACT	ERIS	TTCS	-						
			-					6 am			s					
								no a			~					
								lin								
20			(xi)					PTIO		FO T	ח או	. 26	7.			
			,,,,	DDQ	OLIV.		00111		м. Б	בע ד	ט אי	. 20	٠.			
	Glu	Glv	I.eu	Δτα	T.eu	T.A.11	I.011	Ser	T.eu	Dro	בו מ	λla	Len	Dro	λ~~	C
	1	,		9	5	204		501	<u> </u>	10	nau	niu	Leu	FIO	15	361
	_				,					10					13	
25	Cve	Cve	Hie	Dro	2~~	Trans.	T and	Pro	Val	Vaa						
	0,0	0,0	*****	20	n. g	11p	nea	FIO	25	naa						
				20					23					•		
30	(2)	TNE	ORMA	TTON	EOB	SEO.	י חד	NO:	268.							
	(4,		O1441	11011		SEQ	10,		200.							
			(i)	SEOU	FNCE	СНУ	יוייטעק	ERIS	TCC							
			( - /					21 a			de .					
								no a		acı	us		•			
35								lin								
			(vi)					PTIO		EO T	ח אים	. 26	٥.			
			(242)	JUQ	OLIVC.		JUNI	Ļ I I O.	IV. 3.	EQ I	D NO	. 20	٥.			
	Met	Phe	His	Glv	Tla	Pro	Δĺa	Thr	Dro	Gliv	Tle	Gly	λl =	Dro	Gly	y c.n.
	1		****	013	5		714	****	110	10	110	Gry	AIG	FIU	15	MS11
40	•				,					10					13	
	Tage	Pro	Glu	Leu	The con-	Gliv	Glu.	Val	T 140	T 011	<b>Тъ със</b>	T	λ a.m.	71.	3	G1
	D <sub>2</sub> O	110	Giu	20	ıyı	Giu	GIU	vai		Tea	ıyı	гуs	MSII		Arg	GIU
				20					25					30		
	<b>7~~</b>	C1	7	(Th	3	3	<b>V</b>		<b>~</b> 1	•	Db -		1			_,
45	Arg	GIU		ıyr	ASP	ASI	met	Ala	GIU	Leu	Pne	Ala			Lys	Thr
+5			35					40					45			
				_												
	Met		Ala	Leu	Glu	Lys		Tyr	Ile	Lys	Asp		Val	Ser	Pro	Ser
		50					55					60				
50	_														•	
50	Glu	Tyr	Thr	Ala	Ala	Cys	Ser	Arg	Leu	Leu	Val	Gln	Tyr	Lys	Ala	Ala
	65					70					75					80
	Phe	Arg	Gln	Val	Gln	Gly	Ser	Glu	Ile	Ser	Ser	Ile	Àsp	Glu	Phe	Cys
					85					90					95	
55																
	Arg	Lys	Phe	Arg	Leu	Asp	Cys	Pro	Leu	Ala	Met	Glu	Arg	Ile	Lys	Glu
				100		-	-		105				_	110	-	
	Asp	Arg	Pro	Ile	Thr	Ile	Lys	Asp	Asp	Lys	Gly	Asn	Leu	Asn	Ara	Cvs
50	_	_	115				•	120	-	- :-	-		125		3	

	Ile Ala Asp Val Val Ser Leu Phe Ile Thr Val Met Asp Lys Leu Arg 130 135 140
5	Leu Glu Ile Arg Ala Met Asp Glu Ile Gln Pro Asp Leu Arg Glu Leu 145 150 155 160
10	Met Glu Thr Met His Arg Met Ser His Leu Pro Pro Asp Phe Glu Gly 165 170 175
	Arg Gln Thr Val Ser Gln Trp Leu Gln Thr Leu Ser Gly Met Ser Ala 180 185 190
15	Ser Asp Glu Leu Asp Asp Ser Gln Val Arg Gln Met Leu Phe Asp Leu 195 200 205
	Glu Ser Ala Tyr Asn Ala Phe Asn Arg Phe Leu His Ala 210 215 220
20	
	(2) INFORMATION FOR SEQ ID NO: 269:
25	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 3 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 269:</li> </ul>
30	Nob Tue Ve
50	Met Lys Xaa 1
	·
35	(2) INFORMATION FOR SEQ ID NO: 270:
	(i) SEQUENCE CHARACTERISTICS:
	<ul><li>(A) LENGTH: 49 amino acids</li><li>(B) TYPE: amino acid</li></ul>
40	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 270:
45	Met Gln Ala Pro Phe Xaa His Phe Ser Phe Arg Met Phe Ser Asn Lev 1 5 10 . 15
43 <u>.</u>	Tyr Cys Phe Ser Asp Phe Gln Pro Asn Ile Ser Pro Cys Pro Leu Cys 20 25 30
50	His Cys Ile Leu Pro Xaa His His His Val Phe Leu Leu Leu Ala Val 35 40 45
	Xaa
55	
	(2) INFORMATION FOR SEQ ID NO: 271:
	(i) SEQUENCE CHARACTERISTICS:
60	(A) LENGTH: 52 amino acids

(A) LENGTH: 52 amino acids

	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 271:
5	Met Lys Leu Val Thr Met Phe Asp Lys Leu Ser Arg Asn Arg Val II.  1 5 10 15
10	Gln Pro Met Gly Met Ser Pro Arg Gly His Leu Thr Ser Leu Gln Ass
	Ala Met Cys Glu Thr Met Glu Gln Gln Leu Ser Ser Asp Pro Asp Ser 35 40 45
15	Asp Pro Asp Xaa 50
20	(2) INFORMATION FOR SEQ ID NO: 272:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 32 amino acids
25	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 272:
	Met Ala Val Gly Glu Ala Val Phe Val Pro Leu Gln His Pro Pro Le 1 5 10 15
30	Leu His Gly Ser Pro Ile Pro Lys Leu Leu Pro Gly Pro Leu Leu Xa 20 25 30
35	
40	(2) INFORMATION FOR SEQ ID NO: 273:
40	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 57 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 273:  Met Asn Gly Cys His Arg Arg Lys Arg Leu His Leu Cys Lys Thr II  1 5 10 15
50	Tyr Leu Leu Trp Phe Val Phe Ser Phe Leu Leu Ser Asn Glu Val Va 20 25 30
	Ser Ser His Trp His Ile Leu Arg Ala Val Gln Ile Ile Cys Thr Le 35 40 45
55	Phe His Arg Xaa Ile Ser Ala Phe Xaa 50 55
60	(2) INFORMATION FOR SEQ ID NO: 274:

(2) INFORMATION FOR SEQ ID NO: 274:

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 22 amino acids
                    (B) TYPE: amino acid
 5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 274:
      Met Gly Trp Val Ser Ser Pro His Val Lys Arg Arg Glu Cys Val Leu
                                           10
10
     Lys Lys Pro Phe Phe Xaa
                   20
15
      (2) INFORMATION FOR SEQ ID NO: 275:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 275:
      Met Phe Asn Phe Phe Lys Asn Pro Leu Leu Thr Cys Leu Phe Ile Ser
25
                                           10
      Cys Tyr Leu Tyr Leu Ser Leu Leu Val Asn Lys Val Leu Phe Ala Glu
                   20
30
      Glu Gly Leu Cys Cys Thr Tyr Cys Thr Thr Ser Asn Thr Gly Glu Gly
                                   40
      Gly Val Xaa
           50
35
      (2) INFORMATION FOR SEQ ID NO: 276:
40.
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 2 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 276:
45
      Met Xaa
       1
50
      (2) INFORMATION FOR SEQ ID NO: 277:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 66 amino acids
55
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 277:
      Met Leu Cys Thr Ile Leu Thr Val Val Ile Ile Ile Ala Ala Gln Thr
60
                        5
                                          10
```

	Thr	Arg	Thr	Thr 20	Gly	Ile	Pro	Lys	Asn 25	Ala	Pro	Gly	Pro	Ala 30	Pro	Leu
5	Суѕ	Ala	Pro 35	Arg	Ser	Pro	Arg	Leu 40	Phe	Leu	Gln	Xaa	Тут 45	Arg	Gly	Pro
0	Asn	Gly 50	Arg	Pro	Ala	His	Pro 55	Phe	Leu	Gly	Pro	Ser 60	Asp	Leu	Asp	Thr
	Ser 65	Xaa									*					
15	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	NO: 2	278:						•	
20			(i) :	() ()	A) L: B) T D) T	ENGT YPE : OPOL	H: 2 ami OGY:	57 a no a lin	mino cid ear	aci						
			(xi)	SEQ	JENCI	E DES	SCRI	PTIO	N: S	EQ II	ON C	: 278	3:			
25	Met 1	Leu	Gly	Ala	Lys 5	Pro	His	Trp	Leu	Pro 10	Gly	Pro	Leu	His	Ser 15	Pro
	Gly	Leu	Pro	Leu 20	Val	Leu	Val	Leu	Leu 25	Ala	Leu	Gly	Ala	Gly .30	Trp	Ala
30	Gln	Glu	Gly 35	Ser	Glu	Pro	Val	Leu 40	Leu	Glu	Gly	Glu	Cys 45	Leu	Val	Val
35	Cys	Glu 50	Pro	Gly	Arg	Ala	Ala 55	Ala	Gly	Gly	Pro	Gly 60	Gly	Ala	Ala	Leu
_	Gly 65	Glu	Ala	Pro	Pro	Gly 70	Arg	Val	Ala	Phe	Xaa 75	Ala	Val	Arg	Ser	His 80
10	His	His	Glu	Pro	Ala 85	Gly	Glu	Thr	Gly	Asn 90	Gly	Thr	Ser	Gly	Ala 95	Ile
	Tyr	Phe	Asp	Gln 100	Val	Leu	Val	Asn	Glu 105	Gly	Gly	Gly	Phe	Asp 110	Arg	Ala
15	Ser	Gly	Ser 115	Phe	Val	Ala	Pro	Val 120	Arg	Gly	Val	Tyr	Ser 125	Phe	Arg	Phe
	His	Val 130	Val	Lys	Val	Tyr	Asn 135	Arg	Gln	Thr	Val	Gln 140	Val	Ser	Leu	Met
	Leu 145	Asn	Thr	Trp	Pro	Val 150	Ile	Ser	Ala	Phe	Ala 155	Asn	Asp	Pro	Asp	Val 160
55	Thr	Arg	Glu	Ala	Ala 165	Thr	Ser	Ser	Val	Leu 170	Leu	Pro	Leu	Asp	Pro 175	Gly
	Asp	Arg	Val	Ser 180	Leu	Arg	Leu	Arg	Arg 185	Gly	Xaa	Ser	Thr	Gly 190	Trp	Leu
50	Glu	Tle	ī.eu	Lve	Dha	Len	محس	וום. ז	Dro	Hic	ĭ.en	Pro	Sar	ĭ.e.ı	Lve	Δen

WO 98/54963

			195					200					205				
5	Pro	Ser 210	Leu	Ser	Ser	Thr	Arg 215	Ile	Gln	Pro	Leu	Thr 220	Thr	Phe	Phe	Cys	
5	Pro 225	Leu	Leu	Pro	Хаа	Lys 231	Glm	Уаа	Lys	Gln	Xaa 235	Хаа	Xaa	Ser	Leu	Trp 240	
10	Leu	Leu	Ser	His	Leu 245	Phe	Ala	Trp	Glu	Pro 250	Val	Pro	Asn	Thr	Gln 255	Val	
	Xaa																
15																	
	(2)	⊐NF	OFMAC	TON	FCP.	SEÇ	ID I	NG: 2	279:								
20			(i) : (xi)	()	A) I B) T D) T	enge YPE: OPCL	H: 1 ami CGY:	.03 a no a lin	mino cid ear	aci		: 27	9:				
25	Met 1		<b>P</b> ro	Arg	Ala 5	Leu	Pro	Gly	Ser	Ala 10	Val	Leu	Ala	Ala	Ala 15	Val	
30	Phe	∵al	Gly	Gly 20	Ala	Val	Ser	Ser	Pro 25	Leu	Val	Ala	Pro	Asp 30	Asn	Gly	
	Ser	Ser	Arg 35	Thr	Leu	His	Ser	73 YEG	Thr	Glu	Thr	Thr	Pro 45	Ser	Pro	Ser	
35	Asn	Asp 50	Thr	Gly	Asn	Gly	His 55	Pro	Glu	Tyr	Ile	Ala 60	Tyr	Ala	Leu	Val	
	Pro 65		Phe	Phe	Ile	Met 70	Gŀy	Leu	Phe	Gly	Val 75	Leu	Ile	Xaa	Pro	Xaa 80	
40	Xaa	ĭaa	Lys	Lys	Lys 85	Gl <sub>?</sub>	Tyr	Arg	Cys	Thr 90	Thr	Glu	Ala	Glu	Gln 95	Asp	
45	Ile	Glu	Glu	Glu 100	Lys	G].	Xaa										
•	(2)	INF	CRMA	TICN	FCR	ಽಪಾ	ID :	NC:	280:								
50			(i)		ENCE (A) I (B) I	engi Ype :	H: 3	ns 88 ino a	nino cid		is						
55			(xi)		UENC					EQ I	D NC	): 28	: 0				
	Met 1		7al	Thr	Leu 5	Ser	Ser	· ieu	Gly	Phe 10		Val	Leu	Leu	Ser 15	Leu	
60	Leu	Phe	?ro	Trp	_	The	Asp	Gln	Gly		Gly	Pro	Ala	Thr		Tyr	

Xaa

5

(2) INFORMATION FOR SEQ ID NO: 281:

(i) SEQUENCE CHARACTERISTICS:

10 (A) LENGTH: 43 amino

(A) LENGTH: 43 amino acids(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 281:

Met Val Leu Gly Leu Leu Leu Leu Leu Xaa Phe Phe Ser Phe Ser Ser 1 5 10 15

Ser Pro Ser Pro Ser Ser Ser Leu Leu Leu Leu Ser Ser Phe Phe Phe 20 25 30

20
Gln Ser Leu Ala Leu Ser Pro Arg Leu Glu Xaa

35 . 40

25

30

- (2) INFORMATION FOR SEQ ID NO: 282:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 21 amino acids
  - (B) TYPE: amino acid
    - (D) TOPOLOGY: linear
    - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 282:

Glu Trp Leu Val Phe Thr Phe Leu Leu Val Phe Gly Ser Pro Leu Gly 15

Lys Gly Pro Leu Xaa 20

40

45

55

- (2) INFORMATION FOR SEQ ID NO: 283:
- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 70 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 283:

50 Met Ile Arg Ala Leu Ser Leu Phe Leu Leu Ile Phe Asp Ala Ala Leu
1 10 15

Phe Ser Leu Ser Val Phe Val Phe Ile Gly His Leu Leu Pro Met Pro

Lys Gly Thr Gly Leu His Ser Cys Ala Lys His Leu Ile Lys Ser Leu 35 40 45

Lys Glu Asn Val Leu Pro Leu Met Asn Tyr Pro Asp Cys Lys Leu Lys 50 55 60

```
Ile Asn Ile Ser Pro Xaa
 5
      (2) INFORMATION FOR SEQ ID NO: 284:
             (i) SEQUENCE CHARACTERISTICS:
10
                    (A) LENGTH: 75 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 284:
15
     Met Gly Lys Leu Ile Arg Leu Ser Val Met Val Met Ser Val Arg Arg
                       5
       1
                                          10
     Leu Phe Ser Ile Tyr Trp Val Leu Ser Thr Val Pro Asp Ala Val Gly
                                      25
20
      Ser Arg Gly Gly Met Glu Glu Glu Cys Ser Arg Gly Leu Cys Cys Val
                                  40
                                                      45
      Ala Gly Gln His Lys Gln Ala Lys Gly Lys Arg Gln Ala Trp Asn Lys
25
     Gly Gly Glu Tyr Gln Cys Val Thr Tyr Cys Xaa
                          70
30
      (2) INFORMATION FOR SEQ ID NO: 285:
             (i) SEQUENCE CHARACTERISTICS:
35
                    (A) LENGTH: 33 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 285:
40
     Met Pro Ala Leu Val Thr Leu Leu Leu Phe Pro Leu Leu Pro Leu
                      5
                                10
     Met Glu Ala Ser Cys His Val Met Arg Cys Pro Met Glu Arg Pro Thr
                  20
                                      25
45
     Xaa
50
      (2) INFORMATION FOR SEQ ID NO: 286:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 17 amino acids
55
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 286:
     Glu Ala Pro Trp Gly Leu Leu Lys Leu Leu Leu Leu Leu Ala Val Phe
60
                                          10
```

Xaa 5 (2) INFORMATION FOR SEQ ID NO: 287: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 17 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 287: 15 Met Gln Gln Lys Gln Lys Lys Ala Asn Glu Lys Lys Glu Glu Pro Lys 5 10 Xaa 20 (2) INFORMATION FOR SEQ ID NO: 288: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 38 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 288: 30 Met Gln Arg Lys Val Ser Asp Phe Ile Ile His Gln Arg Leu Thr Val 5 Asn Leu Cys Val Ile Ser Phe Phe Phe Phe Leu Pro Ile Cys Ile Phe 35 25 Ser Leu Ala Lys Lys Xaa 35 40 (2) INFORMATION FOR SEQ ID NO: 289: (i) SEQUENCE CHARACTERISTICS: 45 (A) LENGTH: 12 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 289: 50 Met Ala Leu Leu Ile Ser Ser Leu Ile Trp Ser Xaa 1 5

55 (2) INFORMATION FOR SEQ ID NO: 290:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 35 amino acids

(B) TYPE: amino acid

60 (D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 290:													
5.	Met Gln Met Phe Thr Val Ser Leu Leu Leu Ser Leu Leu Leu Arg Ser 1 5 10 .15													
	Thr Asp Gln Asn His Leu Gln Leu Leu Val Gly Arg Glu Asp His Tyr 20 25 30													
10	Gly Gly Xaa 35													
15	(2) INFORMATION FOR SEQ ID NO: 291:													
20	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 15 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 291:</li> </ul>													
	Met Ser Glu Ser Ala Cys Ile Leu Asn Asn Gln Lys Glu Leu Xaa 1 5 10 15													
25														
	(2) INFORMATION FOR SEQ ID NO: 292:													
30	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 44 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>													
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 292:													
35	Met Asp Leu Asp Arg Val Lys Ala Glu Ala Thr Glu Asp Ile Thr Ser 1 5 10 15													
40	Gly Val Leu Cys Leu Leu Phe Leu Arg Leu Pro Pro Asn Ser Cys Ile 20 25 30													
	Phe Pro Ser Ala Val Leu Gly Ser Thr Arg Thr Xaa 35 40													
45	(2) INFORMATION FOR SEQ ID NO: 293:													
50	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 136 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 293:</li> </ul>													
55	Val Val Gly Thr Gly Thr Ser Leu Ala Leu Ser Ser Leu Leu Ser Leu  1 5 10 15													
	Leu Leu Phe Ala Gly Met Gln Met Tyr Ser Arg Gln Leu Ala Ser Thr 20 25 30													

Glu Trp Leu Thr Ile Gln Gly Gly Leu Leu Gly Ser Gly Leu Phe Val

PCT/US98/11422

			35					40					45			
5	Phe	Ser 50	Leu	Thr	Ala	Phe	Asn 55	Asn	Leu	Glu	Asn	Leu 60	Val	Phe	Gly	Lys
3	Gly 65	Phe	Gln	Ala	Lys	Ile 70	Phe	Pro	Glu	Ile	Leu 75	Leu	Cys	Leu	Leu	Leu 80
10	Ala	Leu	Phe	Ala	Ser 85	Gly	Leu	Ile	His	Arg 90	Val	Cys	Val	Thr	Thr 95	Суз
	Phe	Ile	Phe	Ser 100	Met	Val	Gly	Leu	Tyr 105	Tyr	Ile	Asn	Lys	Ile 110	Ser	Ser
15	Thr	Leu	Tyr 115	Gln	Ala	Ala	Ala	Pro 120	Val	Leu	Thr	Pro	Ala 125	Lys	Val	Thr
20	Gly	Lys 130	Ser	Lys	Lys	Arg	Asn 135	Xaa								
	(2)	INF	ORMA!	NOI	FOR	SEQ	ID I	VO: 2	294:							
25			(i) :	(	A) L B) T	ENGT YPE:	H: 3 ami	4 am no a	ino cid	: acid	s					
30			(xi)				OGY: SCRI			EQ I	D NO	: 29	4 :			•
	Met 1	Phe	Ile	Phe	Leu 5	Phe	Leu	Cys	Val	Leu 10	Ser	Arg	Lys	Ile	Gln 15	Glu
35	Glu	Tyr	Tyr	Arg 20	Leu	Phe	Lys	Asn	Val 25	Pro	Cys	Cys	Phe	Gly 30	Cys	Leu
	Arg	Xaa													·	
40																
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: 2	295 :							
45				(	A) L B) T D) T	ENGT YPE : OPOL	ami OGY:	37 a no a lin	mino cid ear	aci		: 29	5:			
50	Met 1	Arg		_										Leu	Ala 15	Gly
	Ala	Pro	Ala	Ala 20	Arg	Pro	Thr	Pro	Pro 25	Thr	Cys	Tyr	Ser	Arg 30	Met	Arg
55	Ala	Leu	Ser 35	Gln	Glu	Ile	Thr	Arg 40	Asp	Phe	Asn	Leu	Leu 45	Gln	Val	Ser
60	Glu	Pro 50	Ser	Glu	Pro	Cys	Val 55	Arg	Тут	Leu	Pro	Arg 60	Leu	Tyr	Leu	Asp

(2) INFORMATION FOR SEQ ID NO: 298:

	Ile 65	His	Asn	Tyr	Cys	Val 70	Leu	Asp	Lys	Leu	Arg 75	Asp	Phe	Val	Ala	Ser 80
5	Pro	Pro	Cys	Trp	Lys 85	Val	Ala	Gln	Val	Asp 90	Ser	Leu	Lys	Asp	Lys 95	Ala
10	Arg	Lys	Leu	туr 100	Thr	Ile	Met	Asn	Ser 105	Phe	Cys	Arg	Arg	Asp 110	Leu	Val
10	Phe	Leu	Leu 115	Asp	Asp	Cys	Asn	Ala 120	Leu	Glu	Tyr	Pro	Ile 125	Pro	Val	Thr
15	Thr	Val 130	Leu	Pro	Asp	Arg	Gln 135	Arg	Xaa	-	-			٠		
20	(2)	INF		TION SEQU	ENCE	CHA	RACT	ERIS			s					
25			(xi)		D) T	YPE: OPOL E DE	OGY:	lin	ear	EO I	D NO	: 29	6:			
	Met 1	Trp		Leu						•				Xaa	Leu 15	Val
30	Leu	Leu	Phe	Pro 20	Arg	Gly	Trp	Ser	Gln 25	Pro	Gly	Thr	His	Lys 30	Arg	Gln
35	Ile	Leu	Val 35	Asn	Xaa	Ala	Ser	Leu 40	Pro	Gly	Gly	Cys	Leu 45	Leu	Pro	Trp
33	Ile	Trp 50		Gly	Ala	Ala	Leu 55	Arg	Phe	Xaa			-			
40	(2)	INF	ORMA	TION	FOR	SEQ	ID:	NO:	297 :							
45				(	A) I B) T D) T	ENGI YPE : OPOL	H: 3 ami OGY:	5 an no a lir	nino Icid Iear	acid		): <b>2</b> 9	7:			
50	Met 1		Arg	Arg	Ala 5		Ala	Ser	Ile	Phe 10	Val	Leu	Pro	Lys	Thr 15	
	Leu	Phe	· Val	Leu 20		Pro	Ala	Phe	Pro 25		Pro	Ala	Val	Gly 30		Pro
55	Val	Pro	Xaa 35													
	٠.												•			

5			(xi)	(	A) L B) T D) T UENC	YPE: OPOL	ami OGY:	no a lin	cid ear			: 29	8:			
10	Ser 1	Cys	Tyr	Ile	Thr 5	Pro	Trp	Ser	Lys	Ile 10	Gln	Ser	Phe	Ser	Leu 15	Ser
10	Leu	Phe	Gln	Phe 20	Ile	Leu	Gln	Glu	Val 25	Asn	Ile	Thr	Leu	Pro 30	Glu	Asn
15	Ser	Val	Trp 35	Tyr	Glu	Arg	Tyr	Lys 40	Phe	Asp	Ile	Pro	Val 45	Phe	His	Leu
	Asn	Gly 50	Gln	Phe	Leu	Met	Met 55	His	Arg	Val	Asn	Thr 60	Ser	Lys	Leu	Glu
20	Lys 65	Gln	Leu	Leu	Lys	Leu 70	Glu	Gln	Gln	Ser	Thr 75	Gly	Xaa	Xaa		
25	(2)	INFO	ORMA?	rion	FOR	SEQ	ID I	vo: 2	299:					•		
30				(	A) L B) T D) T	ENGT YPE: OPOL	H: 9 ami: OGY:	5 am no a lin	ino cid ear	acid		: 29	9:			
35	Met 1	Phe	Val	Leu	Phe 5	Ser	Leu	Pro	Lys	Tyr 10	Ala	Gly	Leu	Arg	Leu 15	Pro
<i></i>	Ile	Pro	Gly	Leu 20	Ser	Ala	Leu	Leu	Val 25	Phe	Leu	Leu	Ser	Leu 30	Phe	Ser
40	Arg	Arg	Ala 35	Gln	Val	Glu	Leu	Thr 40	Thr	Gly	Arg	Glu	Thr 45	Leu	Pro	Lys
	Asn	Leu 50	Gln	Gly	Тут	Phe	Pro 55	Glu	Phe	Gly	Phe	Gln 60	Val	Gln	Asn	Phe
45	Leu 65	Ser	Cys	Lys	Ile	Тут 70	Ala	Ala	Ser	Gln	Lys 75	Gln	Pro	Leu	Pro	Pro 80
50	Leu	Tyr	Gln	Leu	Arg 85	Phe	Tyr	Leu	Lys	His 90	Met	Gly	Leu	Pro	Хаа 95	
	(2)	INF	ORMA!	rion	FOR	SEQ	ID 1	vo: :	300:							
55			(i)	(	ENCE A) L B) T D) T	ENGT YPE :	H: 4 ami	4 am no a	ino cid		s					
60			(xi)	SEQ						EQ I	D NO	: 30	0:			

	met 1	Ser	Ser	His	Trp 5	Thr	Leu	Lys	Ile	Leu 10	Leu	Val	Pro	Leu	Phe 15	Tyr
5	Leu	Ser	Leu	Glu 20	Phe	Pro	Ser	Gly	Phe 25	Val	Leu	Cys	Leu	Ala 30	Asn	Asp
	Leu	Gly	Tyr 35	His	Phe	Ser	Ser	Arg 40	Val	Arg	Ser	Xaa				
10																
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: :	301:			٠.				
15			(i)	(	A) L B) T		H: 3 ami	1 am no a	ino cid	: acid	s					
			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 30	1:		•	
20	Met 1	Leu	Val	Val	Asn 5	Ile	Asn	Leu		Phe 10	Leu	Leu	Phe	Phe	Ile 15	Phe
25	Leu	Cys	Tyr	Leu 20	Asp	Ala	Cys	Ile	Asn 25	Val	Phe	Cys	Phe	Tyr 30	Xaa	
	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	<b>10:</b> 3	302:							
•																
30	,			(	A) L B) T D) T	ENGT YPE: OPOL	H: 1 ami OGY:	13 a no a lin	mino cid ear	aci		. 20	<b>.</b>			
30			(xi)	() () SEQ	A) L B) T D) T UENC:	ENGT YPE: OPOL E DE	H: 1 ami OGY: SCRI	13 a no a lin PTIO	mino cid ear N: S	aci	D NO					
	Met 1	Pro	(xi)	() () SEQ	A) L B) T D) T UENC:	ENGT YPE: OPOL E DE	H: 1 ami OGY: SCRI	13 a no a lin PTIO	mino cid ear N: S	aci	D NO			Leu	Thr 15	Leu
	1		(xi) Val	() () SEQ	A) L B) T D) T UENC: Pro 5	ENGT YPE: OPOL E DE	H: 1 ami OGY: SCRI Arg	13 a no a lin PTIO	mino cid ear N: Si	aci EQ II Ala	D NO Leu	Leu	Ser		15	
35	1 Ala	Phe	(xi) Val Ala	() () () SEQUE Leu Val 20	A) L B) T D) T UENC: Pro 5	ENGT YPE: OPOL E DE Gly Cys	H: 1 ami OGY: SCRI Arg	13 a no a lin PTIO Thr	mino cid ear N: Si Thr Val 25	EQ II	D NO Leu Ala	Leu Gly	Ser Pro	Cys 30	15 Val	Pro
35	1 Ala Arg	Phe Ser	(xi) Val Ala His 35	() () () () () () () () () () () () () (	A) L B) T D) T UENC: Pro 5 Pro	ENGT YPE: OPOL E DE Gly Cys	H: 1 ami OGY: SCRI Arg Ser	13 a no a lin PTIO Thr Gly Trp 40	mino cid ear N: Si Thr Val 25 Glu	EQ II Ala 10 Glu	D NO Leu Ala Ser	Leu Gly Val	Ser Pro Cys 45	Cys 30 Val	15 Val Thr	Pro
335 440	1 Ala Arg	Phe Ser Thr 50	(xi) Val Ala His 35	(()()()()()()()()()()()()()()()()()()(	A) L B) T D) T UENC Pro 5 Pro Cys	ENGT YPE: OPOL E DE Gly Cys Ser	H: 1 ami OGY: SCRI Arg Ser Ser Trp 55	13 a no a lin prior Thr Gly Trp 40	mino cid ear N: S Thr Val 25 Glu Ala	Ala 10 Glu Ala	D NO Leu Ala Ser	Cly Val Leu 60	Ser Pro Cys 45	Cys 30 Val Pro	15 Val Thr	Pro Ser Ala
35 40	Ala Arg Ser Ala 65	Phe Ser Thr 50	(xi) Val Ala His 35 Pro	(()()()()()()()()()()()()()()()()()()(	A) L B) T D) T UENC: Pro 5 Pro Cys Gly Xaa	ENGT YPE:: OPPOL E DE: Gly Cys Ser Ser Ala 70	MH: 1 ami OGY: SCRI Arg Ser Trp 55 Ala	13 a no a lin a li	mino cid ear N: S Thr Val 25 Glu Ala	Ala 10 Glu Ala	D NO Leu Ala Ser Ala Pro 75	Leu Gly Val Leu 60	Ser Pro Cys 45 Phe	Cys 30 Val Pro Gln	15 Val Thr Ser	Pro Ser Ala Gly 80
335 440	Ala Arg Ser Ala 65 Asp	Phe Ser Thr 50 Trp	(xi) Val Ala His 35 Pro His	(()()()()()()()()()()()()()()()()()()(	A) L B) T D) T UENC:  Pro 5 Pro Cys Gly Xaa Gly 85	ENGT YPE: OPOL E DE: Gly Cys Ser Ser Ala 70	H: 1 ami OGY: SCRI Arg Ser Trp 55 Ala Met	13 a no a lin prio Thr Gly Trp 40 Arg Trp	mino cid ear N: S Thr Val 25 Glu Ala Asp	Ala Arg	D NO Leu Ala Ser Ala Pro 75	Leu Gly Val Leu 60 Trp	Ser Pro Cys 45 . Phe Thr	Cys 30 Val Pro Gln	15 Val Thr Ser Thr Gly 95	Pro Ser Ala Gly 80 Gly

WO 98/54963 PCT/US98/11422

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(2) INFORMATION FOR SEQ ID NO: 303:
             (i) SEQUENCE CHARACTERISTICS:
5
                    (A) LENGTH: 14 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 303:
10
     Thr His Ile His Thr His Ile Ile Cys Ser Ser Val Xaa
                        5
                                           10
15
      (2) INFORMATION FOR SEQ ID NO: 304:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 35 amino acids
                    (B) TYPE: amino acid
20
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 304:
     Met Glu Asn Phe Phe Phe Ser Phe Tyr Leu Phe Leu Ile Thr Leu Ile
       1
                        5
25
      Pro Asn Gly Arg Thr Leu Ser Thr Thr Ala Asp His Cys Lys Ile Pro
                                       25
                   20
      Cys Ile Xaa
30
               35
      (2) INFORMATION FOR SEQ ID NO: 305:
35
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 35 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
40
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 305:
     Met Glu Leu Trp Glu Leu Ala Leu Cys Leu Leu Val Ala Leu Ser Ala
45
      His Met Phe Thr Val Gln Leu Leu Ala Asp Leu Gly Phe Leu Phe Gly
                                  - 25
      Gly Phe Xaa
               35
50
      (2) INFORMATION FOR SEQ ID NO: 306:
55
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 82 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 306:
60
```

	Met 1	GIY	Aia	ΞΙΆ	Ile 5	Leu	λla	Leu	Len	Leu 10		Leu	Glu	Ser	Val 15	Leu
5	Thr	Cīvs	Ser	Trp 20	Ile	Ser	Val	Ser	Thr 25	Ser	Glu	Arg	Gln	Leu 30	Trp	Gln
1 .	Ser	Ser	Gln 35	Lys	Ala	Tix	Ile	Leu 40	Ser	Leu	Lys	Leu	Asp 45	Ser	Cys	Phe
10	Суѕ	Gly 50	His	Ser	Gly	Leu	Lys 55	Gly	Lys	Asn	Glu	Asp 60	Thr	Asp	Ser	Ser
15	Val 65	Pro	Ile	Ile	Pro	Ser 70	Lys	Thr	His	Thr	His 75	Leu	Gly	Lys	His	Leu 80
	Ile	Xaa														
20	(2)	INFO	CEMA:	ricn	FCR.	SEQ	ID I	no: 3	307:		-					
25			(i)	€.	A) I	en Gr	H: 7	ERIS 2 am	ino		s					
25			(xi)	(	D) T	CPOL	OGY:	no a lin PTIC	ear	EQ II	ои с	: 30	7:			
30	Met 1	Phe	Tyr	Phe	Val 5	Leu	Phe	Ile	Tyr	Ser 10	Ser	Ser	Glu	Thr	Trp 15	Ser
	Gly	Ser	7al	Ala 20	Gln	Asp	Gly	Val	His 25	Gly	Val	Ile	Ile	Gly 30	His	Cys
35	Ser	Val	Glu 35	Leu	Pro	Gly	Ser	Gly 40	Ąsp	Pro	Pro	Ala	Ser 45	Ala	Xaa	Leu
<del>4</del> 0	Val	Ala 50	Gly	Thr	Ile	Gly	Thr 55	Суs	Pro	Thr	Met	Pro 60	Gly	Phe	Val	Tyr
	Phe 65	Leu	Asn	ązk	Val	Х <u>аа</u> 70	Asn	Xaa								
45	(2)	INFO	OFMA:	rion	FCR	SEQ	ID 1	<b>vo:</b> 3	308:							
50			(i) .	(	A) I B) T	ENGT YPE:	H: 3 ami	ERIS 4 am no a	ino cid		s					
			(xi)					lin PTIC		EQ II	ON C	: 30	8:			
55	Met 1	qzA	Ser	Thr	Leu 5	Arg	Gln	Gly	Arg	Хаа 10	Leu	Ļeu	Thr	Leu	Val 15	Pro
	Ala	Ser	Leu	Phe 20	Ser	Leu	Thr	Leu	Gly 25	Gly	Pro	Gly	Pro	Trp 30	Lys	Asp
60	Pro	Xaa														

5	(2)	INF	ORMA!	NOI	FOR	SEQ	ID 1	ю: 3	109:		•					
10				(	ENCE A) L B) T D) T UENC	ENGT YPE : OPOL	H: 1 ami: OGY:	15 an no a lin	mino cid ear	aci		: 30	9:			
15	Met 1	Gln	Val	Val	Gly 5	Ser	Trp	Pro	Gly	Arg 10	Val	Gly	Val	Val	Gly 15	Leu
	Ala	Phe	Ser	Leu 20	Val	Ile	Pro	Pro	Pro 25	Ala	Ile	Cys	Ile	Ala 30	Gly	Pro
20	Ala	Pro	Gly 35	Leu	Gly	Gly	Gly	Glu 40	Arg	Gln	Gln	Lys	Gly 45	Leu	Gly	Arg
	Gly	Gly 50	Gly	Gly	Leu	Arg	Asn 55	Cys	Pro	Gly	Arg	Val 60	Gly	Met	Ala	Ala
25	Glu 65	Pro	Gly	Ala	Leu	Leu 70	Cys	Leu	Thr	Ser	<b>Ar</b> g 75	Asp	Gly	Ser	Leu	Leu 80
30	Leu	Ser	Cys	Val	Arg 85	Pro	His	His	Val	Ile 90	Lys	Pro	Lys	Gly	Thr 95 <sub>(</sub>	Ala
50	Lys	Lys	Lys	Lys 100	Lys	Lys	Lys	Lys	Lys 105	Lys	Lys	Lys	Lys	Lys 110	Xaa	Xaa
35	Gly	Gly	Хаа 115													
40	(2)	IŃF		SEQU )	FOR ENCE A) L B) T	CHA ENGT	RACT H: 1	ERIS' 08 a	rics mino		ds					
45			(xi)		D) T UENC					EQ I	D NO	: 31	0:			(
	Met 1	Asp	Leu	Pro	Gln 5	Phe	Ile	Тут	Leu	Phe 10	Ile	Phe	Cys	Phe	Cys 15	Cys
50	Leu	Ala	Ile	Val 20	Asn	Asn	Ala	Ser	Ile 25	Asn	Ile	His	Ile	Gln 30	Val	Ser
55	Met	Trp	Leu 35	Tyr	Val	Phe	Ile	Ser 40	Leu	Gly	Tyr	Leu	His 45	Gly	Ser	Arg
	Ile	Leu 50		His	Asn	Ile	Ile 55	Leu	Cys	Leu	Thr	Ser 60	Gln	Arg	Ile	Ala
60	Lys 65		Phe	Phe	Ile	Val		Ala	Ser	Phe	Thr 75		Pro	Pro	Ala	Met 80

Tyr Lys Asp Phe Tyr Phe Ser Ile Ser Leu His Leu Pro Thr Leu Leu 85 90 Phe Xaa Xaa Xaa Phe Val Phe Ser Leu Leu Pro Pro 100 105 10 (2) INFORMATION FOR SEQ ID NO: 311: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 65 amino acids (B) TYPE: amino acid 15 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 311: Met Cys Ser Pro Ser Leu Ser Ser Ser Pro Pro Pro Leu Leu Gln Val 10 20 Phe Phe Phe Phe Phe Phe Ser Pro His Trp Ala Ala Lys Val Val Pro Gln Trp Lys Xaa Arg His Pro Gln Val Ser Ser Gln Leu Leu Cys 25 40 Phe Leu Arg Val Asn Cys Gln Phe Leu Phe Leu Gln Glu Ile Leu Phe 55 60 30 Xaa 65 35 (2) INFORMATION FOR SEQ ID NO: 312: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 50 amino acids (B) TYPE: amino acid 40 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 312: Met Cys Leu Ser Arg Trp Lys Ile Phe Tyr Thr Leu Leu Ile Leu Phe 10 45 Xaa Xaa Phe Ser Ile Thr Ser Glu Xaa Glu Thr Phe Tyr Met Ile Ile Ile His His Asn Pro Thr Gln Ile Thr Ala Ser Cys Ser Phe Thr Phe 50 35 Leu Xaa 50 55 . (2) INFORMATION FOR SEQ ID NO: 313:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 293 amino acids

## (B) TYPE: amino acid

## (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 313:

	(:	xi) SEQ	UENCE I	DESCRI	PTIO	1: SI	EQ II	ONO:	313	3:			
5	Met Glu i	Arg Pro	Asp Tr 5	p Glu	Thr	Ala	Ile 10	Gln	Lys	Pro	Leu	Cys 15	Ser
10	Leu Pro 1	Ala Gly 20		y Asn	Ala	Leu 25	Ala	Ala	Ser	Leu	Asn 30	His	Tyr
	Ala Gly	Tyr Xaa 35	Gln Va	al Thr	Asn 40	Glu	Asp	Leu	Leu	Thr 45	Asn	Cys	Thr
15	Leu Leu 1 50	Leu Cys	Arg Ar	g Leu 55	Leu	Ser	Pro	Met	Asn 60	Leu	Leu	Ser	Leu
,	His Thr A	Ala Ser	_	eu Arg 70	Leu	Phe	Ser	Val 75	Leu	Ser	Leu	Ala	Trp 80
20	Gly Phe	Ile Ala	Asp Va 85	al Asp	Leu	Glu	Ser 90	Glu	Lys	Tyr	Arg	Arg 95	Leu
25	Gly Glu I	Met Arg 100		ır Leu	Gly	Thr 105	Phe	Leu	Arg	Leu	Ala 110	Ala	Leu
	Arg Thr	Tyr Arg 115	Gly A	g Leu	Ala 120	Tyr	Leu	Pro	Val	Gly 125	Arg	Val	Gly
30	Ser Lys 1	Thr Pro	Ala Se	er Pro 135	Val	Val	Val	Gln	Gln 140	Gly	Pro	Val	Asp
•	Ala His 1 145	Leu Val		eu Glu 50	Glu	Pro	Val	Pro 155	Ser	His	Trp	Thr	Val 160
35	Val Pro		165				170	,				175	
40	Leu Gly	Ser Glu 180		ne Ala	Ala	Pro 185	Met	Gly	Arg	Cys	Ala 190	Ala	Gly
		195			200					205			
45	Leu Arg			215					220				
50	Cys Pro		2:	30				235					240
50	Lys Asp	Gly Lys	245	al Phe	Ala	Val	Asp 250	Gly	Glu	Leu	Met	Val 255	Ser
55	Glu Ala	Val Gli 260	_	ln Val	His	Pro 265	Asn	Tyr	Phe	Trp	Met 270	Val	Ser
	Gly Cys	Val Glu 275	ı Pro P	ro Pro	Ser 280	Trp	Lys	Pro	Gln	Gln 285	Met	Pro	
60	Pro Glu 290	Glu Pro	Leu									•	

(2) INFORMATION FOR SEQ ID NO: 316:

5	(2)	INF	ORMA?	rion	EOB	~	ID 1	<b>10:</b> 3								
					FOR	SEQ			314:							
			(i)	(	A) L B) T	ENGT YPE :	H: 6 ami	ERIS 8 am no a	ino cid		s					
10			(xi)					lin PTIO		EQ I	D NO	: 31	4:			
	Met 1	Pro	Leu	Glu	Gly 5	Phe	Суѕ	Leu	Val	Leu 10	Asp	Ile	Gly	Phe	Leu 15	Leu
15	Val	Met	Leu	Ile 20	Ser	Leu	Ala	Ser	Glu 25	Суѕ	Phe	Thr	Thr	Cys 30	Leu	Asp
20	Ser	Phe	Ser 35	Thr	Thr	Glu	Pro	Gly 40	Cys	Lys	Phe	Tyr	Lys 45	Leu	Leu	His
	Ser	Val 50	Ser	Leu	Leu	Asn	Ile 55	Asn	Phe	Asn	Val	Lys 60	Ser	Leu	Leu	Cys
25	Ser 65	His	Ile	Xaa											•	
30	(2)	INF						NO: 3								
			(i)			CHAI ENGT		ERIS			_					
35							ami	no a	cid	aci	ds					
	•		(xi)	(	D) T	OPOL	ami OGY:		cid ear			: 31	5:			
	Met 1	Pro		SEQ	D) T UENC	OPOL E DE	ami OGY: SCRI	no a lin	cid ear N: S	EQ I	D NO			Leu	Val 15	Phe
40	1		Leu	SEQ Gln	D) T UENC Leu 5	OPOL E DE Ser	ami OGY: SCRI Gly	no a lin PTIO	cid ear N: S Tyr	EQ II Trp 10	D NO	Ser	Leu		15	
40 45	1 Leu	Ser	Leu Leu	Gln Gln 20	D) T UENC Leu 5	OPOL E DE Ser Phe	ami OGY: SCRI Gly Pro	no a lin PTIO Gln	cid ear N: S Tyr Ala 25	Trp 10	D NO Ile Ile	Ser Pro	Leu Cys	Ala 30	15 Leu	Thr
	1 Leu Asp	Ser Val	Leu  Gly 35 Leu	Gln Gln 20 Gly	D) T UENC Leu 5 Pro	OPOL E DE Ser Phe Cys	ami OGY: SCRI Gly Pro Val	no a lin PTIO Gln Gln	cid ear N: S Tyr Ala 25 Cys	Trp 10 Ala	D NO Ile Ile	Ser Pro Leu	Leu Cys Leu 45	Ala 30 Asn	15 Leu Cys	Thr Leu
	1 Leu Asp Cys	Ser Val Ile 50	Leu Gly 35 Leu	Gln Gln 20 Gly Phe	D) TUENC	OPOL E DE Ser Phe Cys	ami OGY: SCRI Gly Pro Val Thr 55	no a lin PTIO Gln Gln Ile 40	cid ear N: S: Tyr Ala 25 Cys	Trp 10 Ala His	D NO Ile Ile Leu	Ser Pro Leu Ser 60	Leu Cys Leu 45 His	Ala 30 Asn Val	15 Leu Cys Leu	Thr Leu Leu
45 50	1 Leu Asp Cys	Ser Val Ile 50 Lys	Leu Gly 35 Leu Met	Gln Gln 20 Gly Phe	D) TUENC	OPOLLEDE DE Ser Phe Cys Leu Ser 70	ami OGY: SCRI Gly Pro Val Thr 55	no a lin	cid ear N: S Tyr Ala 25 Cys Pro	Trp 10 Ala His	Ile Ile Leu Pro 75	Ser Pro Leu Ser 60	Leu Cys Leu 45 His	Ala 30 Asn Val Asp	15 Leu Cys Leu Leu	Thr Leu Leu Ser 80
45 50	Leu Asp Cys Ile 65 Asp	Ser Val Ile 50 Lys	Leu Gly 35 Leu Met	(SEQ) Gln Gln 20 Gly Phe Ser	D) TUENC Leu 5 Pro Ser Thr Leu Thr 85	OPOL E DE Ser Phe Cys Leu Ser 70	ami OGY: SCRI Gly Pro Val Thr 55 Val	no a lin	cid ear N: S Tyr Ala 25 Cys Pro Tyr	Trp 10 Ala His Ser Glu Leu	Ile Ile Leu Pro 75	Ser Pro Leu Ser 60	Leu Cys Leu 45 His	Ala 30 Asn Val Asp	15 Leu Cys Leu Leu	Thr Leu Leu Ser 80

5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 71 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 316:
10	Met Trp Gly Cys Ser Gly Leu Gly His Arg Thr Val Ser Phe Leu Leu 1 5 10 15
	Leu Leu Pro Cys Ser Phe Pro Arg Pro Cys Xaa Leu Phe Gly Leu Ile 20 25 30
15	Pro Ile Ser Arg Pro Cys Lys Val Glu Ala Pro Arg Leu Ser Val Pro 35 40 45
	Xaa Leu Ser Cys Ala Ser His Pro Tyr Cys Asn Cys Pro Met Ser Thr 50 55 60
20	Ser Cys Pro Leu Pro Arg Xaa 65 70
25	(2) INFORMATION FOR SEQ ID NO: 317:
30	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 39 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 317:</li> </ul>
35	Met Leu Asn Val Leu Ser Lys Val Gln Gln Leu Val Ser Xaa Leu Gly 1 5 10 15
	Leu Val Thr Phe Leu Leu Asn His Ser Ala Ala Gly Gly Ser Pro Gln 20 25 30
40	His Arg Trp Leu Leu Xaa 35
45	(2) INFORMATION FOR SEQ ID NO: 318:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 72 amino acids  (B) TYPE: amino acid
50	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 318:
	Met Lys Ala Ile Ala Arg Ala Cys Leu Leu Leu Ser Leu Leu Val Leu  1 5 10 15
55	Pro His Val Val Ser Glu His Leu Phe Trp His His Asn Pro Arg His 20 25 30
60	Pro Val Ile Trp Pro Phe Pro Pro Phe His Leu Ile Ser Cys Ser Val

	Ser	Ala 50	Ser	Thr	Trp	His	Leu 55	Gly	Glu	Xaa	Leu	Leu 60	Leu	Leu	Val	Pro
5	Ile 65	Ala	Pro	Ser	Val	Trp 70	Ser	Xaa							•	
10	(2)		ORMAT		٠	-				<b>:</b>						
15			(xi)	0	B) T D) T	YPE: OPOL	ami OGY:	2 am no a lin PTIO	cid ear			: 31	9:		·	
	Met 1	Glu	Gln	Gly	Gly 5	Gly	Pro	Arg	Leu	Leu 10	Leu	Leu	Ile	Pro	Gly 15	Leu
20	Leu	His	Asn	Thr 20	Тут	Leu	Ala	Arg	Pro 25		Asp	Phe	Pro	Ala 30	Gln	Gly
25	Thr	Thr	Glu 35	Asn	Thr	Glu	Cys	Gln 40	Gly	Ser	Pro	Ser	Pro 45	Ile	Ser	His
23	Leu	Gly 50	Lys	Val	Arg	Ser	Leu 55	Asp	Ser	Asn	Thr	Gln 60	Ile	Xaa		
30																
50	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: 3	320:							
35	(2)	INF	(i)	SEQUI ) ) )	ENCE A) L B) T D) T	CHAI ENGT YPE: OPOL	RACT H: 2 ami OGY:	NO: CERISON NO: CE	TICS mino cid ear	aci		: 32	0:			
			(i)	SEQUI ( ( SEQI	ENCE A) L B) T D) T UENC	CHAI ENGT YPE: OPOL E DE	RACT H: 2 ami OGY: SCRI	ERIS 86 a no a lin PTIO	TICS mino cid ear N: S	aci EQ I	D NO			Gly	Ser 15	Val
35	Met 1	Pro	(i) :	SEQU ( ( SEQ Leu	ENCE A) L B) T D) T UENC Phe 5	CHAI ENGT YPE: OPOL E DE	RACT H: 2 ami OGY: SCRI Ser	ERIS 86 a no a lin PTIO	TICS mino cid ear N: S	aci EQ I Thr 10	D NO Leu	Phe	Ser	•	15	
35	Met 1 Thr	Pro	(i) (xi) Leu	SEQU ( ( SEQ Leu Gln 20	ENCE A) L B) T D) T UENC Phe 5	CHAI ENGT YPE: OPOL E DE Phe	RACT H: 2 ami OGY: SCRI Ser Met	ERIS 86 a no a lin PTIO Val	TICS mino cid ear N: S Ser Leu 25	aci EQ I Thr 10 Pro	D NO Leu Trp	Phe Thr	Ser	Thr 30	15 Gly	Glu
35 40 45	Met 1 Thr	Pro Leu Val	(i) (xi) Leu Gln Leu	SEQU ( ( SEQ Leu Gln 20	ENCE A) L B) T D) T UENC Phe 5 Arg	CHAI ENGT YPE: OPOL E DE Phe Gly	RACTH: 2 ami OGY: SCRI Ser Met	ERIS'86 a no a lin PTIO	TICS mino cid ear N: S Ser Leu 25	EQ I Thr 10 Pro	D NO Leu Trp Glu	Phe Thr Leu	Ser Gly Ile 45	Thr 30 Leu	15 Gly Glu	Glu Met
35	Met 1 Thr Gln Asn	Pro Leu Val	(i) (xi) Leu Gln Leu 35	SEQUI ( ( ( ( SEQUI Leu 20 Ala	ENCE A) L B) T D) T UENC Phe 5 Arg Leu Val	CHAMPENGTON CHAMPENGTON COMPOLE DE DE Phe Gly Leu Arg	RACTH: 2 ami OGY: SCRI Ser Met Trp Ser 55	ERIS'86 a no a a linn PTIO Val Phe Pro 40	TICS mino cid ear N: S Ser Leu 25	aci EQ I Thr 10 Pro	D NO Leu Trp Glu Gln	Thr Leu Arg	Ser Gly Ile 45 Leu	Thr 30 Leu Gly	15 Gly Glu Gly	Glu Met Leu
35 40 45	Met 1 Thr Gln Asn Asp 65	Pro Leu Val Val Thr	(i) (xi) Leu Gln Leu 35	SEQUI ( ( ( ( SEQUI Leu Gln 20 Ala Ser	ENCE A) L B) T D) T UENC Phe 5 Arg Leu Val	CHAMPERST CHAMPERST COPOLL FOR DE DE DE DE Gly Leu Arg 70	RACT H: 2 ami OGY: SCRI Ser Trp Ser 55	ERIS'86 a no a a linn PTIO Val Phe Pro 40 Thr	TICS mino cid ear N: S Ser Leu 25 Arg Asp	aci EQ I Thr 10 Pro Phe Arg	D NO Leu Trp Glu Gln Tyr 75	Thr Leu Arg 60	Ser Gly Ile 45 Leu Glu	Thr 30 Leu Gly Phe	Gly Glu Gly Ser	Glu Met Leu Ser 80

Ala Ala Glu Phe Ser Ser Arg Lys Glu Gln Leu Val Phe Leu Ile Asn

			115					120					125			
5	Asn	Tyr 130	Asp	Met	Met	Leu	Gly 135	Val	Leu	Met	Glu	Arg 140	Ala	Ala	Asp	Asp
	Ser 145	Lys	Glu	Val	Glu	Ser 150	Phe	Gln	Gln	Leu	Leu 155	Asn	Ala	Arg	Thr	Glr 160
10	Glu	Phe	Ile	Glu	Glu 165	Leu	Leu	Ser	Pro	Pro 170	Phe	Gly	Gly	Leu	Val 175	Ala
	Phe	Val	Lys	Glu 180	Ala	Glu	Ala	Leu	Ile 185	Glu	Arg	Gly	Gln	Ala 190	Glu	Arg
15	Leu	Arg	Gly 195	Glu	Glu	Ala	Arg	Val 200	Thr	Gln	Leu	Ile	Arg 205	Gly	Phe	Gly
20	Ser	Ser 210	Trp	Lys	Ser	Ser	Val 215	Glu	Ser	Leu	Ser	Gln 220	Asp	Val	Met	Arg
	Ser 225	Phe	Thr	Asn	Phe	Arg 230	Asn	Gly	Thr	Ser	Ile 235	Ile	Gln	Gly	Ala	Leu 240
25	Thr	Gln	Leu	Ile	Gln 245	Leu	Tyr	His	Arg	Phe 250	His	Arg	Val	Leu	Ser 255	Gln
	Pro	Gln	Leu	Arg 260	Ala	Leu	Pro	Ala	Arg 265	Ala	Glu	Leu	Ile	Asn 270	Ile	His
80	His	Leu	Met 275	Val	Glu	Leu	Lys	Lys 280	His	Lys	Pro	Asn	Phe 285	Xaa		
35	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	vo: 3	321:							
			(i) :					ERIS 5 am			5					
10		•	(xi)	(	D) T	OPOL	OGY:	no a lin PTIO	ear	EQ II	ONO:	: 32:	l:			
15	Met 1	Phe	Arg	Ala	Leu 5	Arg	Asp	Leu	Leu	Thr 10	His	Tyr	Pro	Gln	Gln 15	Ile
. •	Leu	Leu	Gln	Val 20	Leu	Val	Val	Met	Ту <del>г</del> 25	Gln	Val	Leu	Gln	Val 30	Trp	Glu
60	Leu	Pro	Trp 35	Pro	Glu	Leu	Ile	His 40	Leu	Gln	Gly	Ile	Val 45	Pro	Thr	Asp
	Gln	Leu 50	His	Leu	Lys	Gln	Xaa 55									
55		•														
	(2)	INFO	ORMAT	MOI	FOR	SEQ	ID N	Ю: 3	22:							
60			(i) S					ERIST 9 am:			3					

WO 98/54963 PCT/US98/11422

•								no a								
								lin		na -		201	_			-
	•		(xi)	SEQ	UENC.	E DE	SCRI.	PITO	N: S	EQ I	ON C	: 32	2:			
5	Asp 1	Phe	Val	Pro	Val 5	Leu	Val	Phe	Val	Leu 10	Ile	Lys	Ala	Asn	Pro 15	Pro
10	Cys	Leu	Leu	Ser 20	Thr	Val	Gln	Tyr	Ile 25	Ser	Ser	Phe	Tyr	Ala 30	Ser	Cys
	Leu	Ser	Gly 35	Glu	Glu	Ser	Tyr	Trp 40	Trp	Met	Gln	Phe	Thr 45	Ala	Ala	Val
15	Glu	Phe 50	Ile	Lys	Thr	Ile	Asp 55	Asp	Arg	Lys	Xaa					
20	(2)	INF	ORMA							:						
				(	A) L B) T	ENGT YPE:	H: 1 ami	20 a no a lin	mino cid		ds					
25			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 32	3:			
	Met 1	His	Pro	Ala	Arg 5	Lys	Leu	Leu	Ser	Leu 10	Leu	Phe	Leu	Ile	Leu 15	Met
30	Gly	Thr	Glu	Leu 20	Thr	Gln	Asp	Ser	Ala 25	Ala	Pro	Asp	Ser	Leu 30	Leu	Arg
35	Ser	Ser	Lys 35	Gly	Ser	Thr	Arg	Gly 40	Ser	Leu	Ala	Ala	11e 45	Val	Ile	Trp
	Arg	Gly 50	Lys	Ser	Glu	Ser	Arg 55		Ala	Lys	Thr	Pro 60	Gly	Ile	Phe	Arg
40	Gly 65		Gly	Thr	Leu	Val 70	Leu	Pro	Pro	Thr	His 75	Thr	Pro	Glu	Trp	Leu 80
	Ile	Leu	Pro	Leu	Gly 85	Ile	Thr	Leu	Pro	Leu 90	Gly	Ala	Pro	Glu	Thr 95	Gly
45	Gly	Gly	Asp	Суs 100	Ala	Ala	Glu	Thr	Trp 105	Lys	Gly	Ser	Gln	Arg 110	Ala	Gly
50	Gln	Leu	Cys 115	Ala	Leu	Leu	Ala	Xaa 120						,		
	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO:	324:							
55			(i)	(	A) L B) T	ENGT YPE:	H: 4 ami	ERIS 4 am no a lin	ino cid		s					
			(xi)					PTIO		EQ I	D NO	: 32	4:			

Phe Phe Leu Val Val Phe Ser Leu Ser Phe Xaa Pro Ser Val Leu Thr Ser Pro Val His Xaa Pro His Cys Cys Gln Xaa Asp Xaa Ile Leu Phe 5 Lys Asn Thr Leu Xaa Xaa Phe Xaa Ala Lys Tyr Xaa 10 (2) INFORMATION FOR SEQ ID NO: 325: (i) SEQUENCE CHARACTERISTICS: 15 (A) LENGTH: 59 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 325: 20 Met Phe Ser Arg Thr Ser Asn Phe Trp Thr Phe Phe Phe Gln Phe Leu 10 Ile Phe Lys Val Phe Leu Val Leu Lys Asn Xaa Phe Thr Ser Gln Lys 25 Ile Xaa Xaa Ile Xaa Xaa Glu Lys Pro Lys Lys Lys Xaa Arg Gly 40 Gly Arg Ala Pro Ser Pro Gln Gly Gly Pro Xaa 30 . 55 50 (2) INFORMATION FOR SEQ ID NO: 326: 35 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 326: Met Gly Leu Leu Ile Phe Met Leu Leu Ile Gly Ile His Ser Gln Cys . 10 45 Ser Xaa 50 (2) INFORMATION FOR SEQ ID NO: 327: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 87 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 327: Met Val Leu Phe Cys Phe Val Leu Phe Cys Phe Val Phe Glu Met Asp. 5 10 60

	Ser	Ser	Ser	Val 20	Thr	Gln	Ala	Gly	Val 25	Gln	Trp	Cys	Asp	Leu 30	Gly	Ser
5	Leu	Gln	Ala 35	Pro	Pro	Pro	Gly	Phe 40	Ser	Pro	Phe	Ser	Cys 45	Leu	Ser	Leu
	Pro	Ser 50	Ser	Trp	Asp	Tyr	Arg 55	Arg	Pro	Pro	Pro	Arg 60	Pro	Ala	Asn	Phe
0	Leu 65	Tyr	Phe	Leu	Val	Glu 70	Thr	Gly	Phe	His	His 75	Val	Ser	Gln	Asp	80
15	Leu	Asp	Leu	Leu	Thr 85	Ser	Xaa								,	
	(2)	INF	ORMA	rion	FOR	SEQ	ID 1	vo: 3	328:							
20				(	A) L B) T D) T	ENGT YPE : OPOL	H: 5 ami: OGY:	38 a no a lin	mino cid ear	aci						
25			(xi)	SEQ	JENC!	E DES	SCRI	PTIO	N: S	EQ II	ON C	: 32	8: .		•	
	Met 1	Ser	Thr	Lys	Lys 5	Leu	Cys	Ile	Val	Gly 10	Gly	Ile	Leu	Leu	Val 15	Phe
30	Gln	Ile	Ile	Ala 20	Phe	Leu	Val	Gly	Gly 25	Leu	Ile	Ala	Pro	Gly 30	Pro	Thr
	Thr	Ala	Val 35	Ser	Tyr	Met	Ser	Val 40	Lys	Cys	Val	Asp	Ala 45	Arg	Lys	Asn
35	His	His 50	Lys	Thr	Lys	Trp	Phe 55	Val	Pro	Trp	Gly	Pro 60	Àsn	His	Cys	Asp
<b>1</b> 0	Lys 65	Ile	Arg	Asp	Ile	Glu 70	Glu	Ala	Ile	Pro	Arg 75	Glu	Ile	Glu	Ala	Asn 80
<del>1</del> 0	Asp	Ile	Val	Phe	Ser 85	Val	His	Ile	Pro	Leu 90	Pro	His	Met	Glu	Met 95	Ser
<b>1</b> 5	Pro	Trp	Phe	Gln 100	Phe	Met	Leu	Phe	Ile 105	Leu	Gln	Leu	Asp	Ile 110	Ala	Phe
	Lys	Leu	Asn 115	Asn	Gln	Ile	Arg	Glu 120	Asn	Ala	Glu	Val	Ser 125	Met	Asp	Val
50	Ser	Leu 130	Ala	Tyr	Arg	Asp	Asp 135	Ala	Phe	Ala	Glu	Trp 140	Thr	Glu	Met	Ala
55	His 145	Glu	Arg	Val	Pro	Arg 150	Lys	Leu	Lys	Cys	Thr 155	Phe	Thr	Ser	Pro	Lys 160
,,,	Thr	Pro	Glu	His	Glu 165	Gly	Arg	Tyr	Tyr	Glu 170	Cys	Asp	Val	Leu	Pro 175	Phe
50	Met	Glu	Ile	Gly		Val	Ala	His	Lys	Phe	Tyr	Leu	Leu	Asn		Arg

	Leu	Pro	Val 195	Asn	Glu	Lys	Lys	Lys 200	Ile	Asn	Val	Gly	11e 205	Gly	Glu	Ile
5	Lys	Asp 210	Ile	Arg	Leu	Val	Gly 215	Ile	His	Gln	Asn	Gly 220	Gly	Phe	Thr	Ļys
10	Val 225	Trp	Phe	Ala	Met	Lys 230	Thr	Phe	Leu	Thr	Pro 235	Ser	Ile	Phe	Ile	Ile 240
	Met	Val	Trp	Tyr	Trp 245	Arg	Arg	Ile	Thr	Met 250	Met	Ser	Arg	Pro	Pro 255	Val
15	Leu	Leu	Glu	Lys 260	Val	Ile	Phe	Aļa	Leu 265	Gly	Ile	Ser	Met	Thr 270	Phe	Ile
	Asn	Ile	Pro 275	Val	Glu	Trp	Phe	Ser 280	Ile	Gly	Phe	Asp	Trp 285	Thr	Trp	Met
20	Leu	Leu 290	Phe	Gly	Asp	Ile	Arg 295	Gln	Gly	Ile	Phe	Tyr 300	Ala	Met	Leu	Leu
25	Ser 305	Phe	Trp	Ile	Ile	Phe 310	Cys	Gly	Glu	His	Met 315	Met	Asp	Gln	His	Glu 320
	Arg	Asn	His	Ile	Ala 325	Gly	Tyr	Trp	Lys	Gln 330	Val	Gly	Pro	Ile	Ala 335	Val
30	Gly	Ser	Phe	Cys 340	Leu	Phe	Ile	Phe	Asp 345	Met	Суѕ	Glu	Arg	Gly 350	Val	Gln
	Leu	Thr	Asn 355	Pro	Phe	Tyr	Ser	Ile 360	Trp	Thr	Thr	Asp	Ile 365	Gly	Thr	Glu
35	Leu	Ala 370	Met	Ala	Phe	Ile	Ile 375	Val	Ala	Gly	Ile	380	Leu	·Cys	Leu	Tyr
40	Phe 385	Leu	Phe	Leu	Cys	Phe 390	Met	Val	Phe	Gln	Val 395	Phe	Arg	Asn	Ile	Ser 400
	Gly	Lys	Gln	Ser	Ser 405	Leu	Pro	Ala	Met	Ser 410	Lys	Val	Arg	Arg	Leu 415	His
45	Tyr	Glu	Gly	Leu 420	Ile	Phe	Arg	Phe	Lys 425	Phe	Leu	Met	Leu	Ile 430	Thr	Leu
	Ala	Cys	Ala 435	Ala	Met	Thr	Val	Ile 440	Phe	Phe	Ile	Val	Ser 445	Gln	Val	Thr
50	Glu	Gly 450	His	Trp	Lys	Trp	Gly 455	Gly	Val	Thr	Val	Gln 460	Val	Asn	Ser	Ala
55	Phe 465	Phe	Thr	Gly	Ile	Tyr 470	Gly	Met	Trp	Asn	Leu 475	Tyr	Val	Phe	Ala	Leu 480
	Met	Phe	Leu	Tyr	Ala 485	Pro	Ser	His	Lys	Asn 490	Tyr	Gly	Glu	Asp	Gln 495	Ser
60	Asn	Gly	Met	Gln 500	Leu	Pro	Cys	Lys	Ser 505	Arg	Glu	Asp	Cys	Ala 510	Leu	Phe

	vai	. Jer	515		. IYL	GIII	GIU	520		: Ser	ATS	ser	525	_	Ser	Phe
5	Ile	530		Asn	Ala	Ala	Ser 535	Gly	Ile	Xaa						
10	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	329:							
15				(	(A) I (B) T (D) T	ENGT YPE : YPOL	H: 2 ami OGY:	ERIS 202 a no a lin PTIO	mino cid ear	aci		: 32	9:			
20	Met 1	Gly	Ile	Ala	Leu 5	Ala	Val	Leu	Gly	Trp 10	Leu	Ala	Val	Met	Leu 15	Cys
	Cys	Ala	Leu	Pro 20	Met	Trp	Arg	Val	Thr 25	Ala	Phe	Ile	Gly	Ser 30	Asn	Ile
25	Val	Thr	Ser 35	Gln	Thr	Ile	Trp	Glu 40	Gly	Leu	Trp	Met	Asn 45	Cys	Val	Val
	Gln	Ser 50	Thr	Gly	Gln	Met	Gln 55	Cys	Lys	Val	Tyr	Asp 60	Ser	Leu	Leu	Ala
30	Leu 65	Pro	Gln	Asp	Leu	Gln 70	Ala	Ala	Arg	Ala	Leu 75	Val	Ile	Ile	Ser	Ile 80
35	Ile	Val	Ala	Ala	Leu 85	Gly	Val	Leu	Leu	Ser 90	Val	Val	Gly	Gly	Lys 95	Cys
,	Thr	Asn	Cys	Leu 100	Glu	Asp	Glu	Ser	Ala 105	Lys	Ala	Lys	Thr	Met 110	Ile	Val
40	Ala	Gly	Val 115	Val	Phe	Leu	Leu	Ala 120	Gly	Leu	Met	Val	Ile 125	Val	Pro	Val
	Ser	Trp 130	Thr	Ala	His	Asn	Ile 135	Ile	Gln	Asp	Phe	Tyr 140	Asn	Pro	Leu	Val
45	Ala 145	Ser	Gly	Gln	Lys	Arg 150	Glu	Met	Gly	Ala	Ser 155	Leu	Tyr	Val	Gly	Trp 160
	Ala	Ala	Ser	Gly	Leu 165	Leu	Leu	Leu	Gly	Gly 170	Gly	Leu	Leu	Cys	Cys 175	Asn

Cys Pro Pro Arg Thr Asp Lys Pro Tyr Ser Ala Lys Tyr Ser Ala Ala

185

190

(2) INFORMATION FOR SEQ ID NO: 330:

Arg Ser Ala Ala Ala Ser Asn Tyr Val Xaa 195 200

180

55

(i) SEQUENCE CHARACTERISTICS:

				(	B) T	YPE:	H: 2 ami	no a	cid	aci	ds					
5			(xi)				OGY: SCRI			EQ I	D NO	: 33	0:			
	Met 1	Ala	Thr	Val	Thr 5	Ala	Thr	Thr	Lys	Val 10	Pro	Glu	Ile	Arg	Asp 15	Va
10	Thr	Arg	Ile	Glu 20	Arg	Ile	Gly	Ala	His 25	Ser	His	Ile	Arg	Gly 30	Leu	Gl
15	Leu	Asp	Asp 35	Ala	Leu	Glu	Pro	Arg 40	Gln	Ala	Ser	Gln	Gly 45	Met	Val	Gl
	Gln	Leu 50	Ala	Ala	Arg	Arg	Ala 55	Ala	Gly	Val	Val	Leu 60	Glu	Met	Ile	Ar
20	Glu 65	Gly	Lys	Ile	Ala	Gly 70	Arg	Ala	Val	Leu	Ile 75	Ala	Gly	Gln	Pro	G1;
	Thr	Gly	Lys	Thr	Ala 85	Ile	Ala	Met	Gly	Met 90	Ala	Gln	Ala	Leu	Gly 95	Pr
25	Asp	Thr	Pro	Phe 100	Thr	Ala	Ile	Ala	Gly 105	Ser	Glu	Ile	Phe	Ser 110	Leu	Gl
30	Met	Ser	Lys 115		Glu	Ala	Leu	Thr 120	Gln	Ala	Phe	Arg	Arg 125	Ser	Ile	G1
	Val	Arg 130	Ile	Lys	Glu	Glu	Thr 135	Glu	Ile	Ile	Glu	Gly 140	Glu	Val	Val	Gl
35	Ile 145	Gln	Ile	Asp	Arg	Pro 150	Ala	Thr	Gly	Thr	Gly 155	Ser	Lys	Val	Gly	Ly:
	Leu	Thr	Leu	Lys	Thr 165	Thr	Glu	Met	Glu	Thr 170	Ile	Tyr	Asp	Leu	Gly 175	Th
10	Lys	Met	Ile	Xaa 180	Ser	Leu	Thr	Lys	Asp 185	Lys	Val	Gln	Ala	Gly 190	Asp	Va
15	Ile	Thr	Ile 195	Asp	Lys	Ala	Thr	Gly 200	Lys	Ile	Ser	Lys	Leu 205	Gly	Arg	Se
	Phe	Thr 210	Arg	Ala	Arg	Glu	Leu 215	Arg	Arg	Tyr		Leu 220	Pro	Asp	Gln	Va
50	Arg 225	Ala	Val	Pro	Arg	Trp 230	Gly	Ala	Pro	Glu	Thr 235	Gln	Gly	Gly	Gly	Ala 24
	His	Arg	Val	Pro	Ala 245	Arg	Asp	Arg	Arg	His 250	Gln	Leu	Ser	His	Pro 255	G1
55	Leu	Pro	Gly	Ala 260	Leu	Leu	Arg									

60 (2) INFORMATION FOR SEQ ID NO: 331:

5					JENCE (A) I (B) I (D) I	LENG TYPE TOPOI	TH: : : am: :COGY	260 á ino á : lir	amino acid near	o ac:		o: 33	31:			•
10	Met 1	Leu	Ala	Leu	Leu 5		Leu	Ser	Gln	Ala 10		Asn	lle	Leu	Leu 15	
	Leu	Lys	Gly	Leu 20	Ala	Pro	Ala	Glu	Ile 25		Ala	Val	. Cys	Glu 30	-	Gly
15	Asn	Phe	Asn 35	Val	Ala	His	Gly	Leu 40		Trp	Ser	Tyr	Tyr 45		Gly	Туз
	Leu	Arg 50		Ile	Leu	Pro	Glu 55		Gln	Ala	Arg	Ile 60		Thr	Tyr	Ası
20	Gln 65	His	Tyr	Asn	Asn	Leu 70	Leu	Arg	Gly	Ala	Val 75		Gln	Arg	Leu	Тут 80
25	Ile	Leu	Leu	Pro	Leu 85	Asp	Cys	Gly	Val	Pro 90	Asp	Asn	Leu	Ser	Met 95	Ala
	Asp	Pro	Asn	Ile 100	Arg	Phe	Leu	Asp	Lys 105	Leu	Pro	Gln	Gln	Thr 110	Gly	Asp
30	Arg	Ala	Gly 115	Ile	Lys	Asp	Arg	Val 120	Tyr	Ser	Asn	Ser	Ile 125	Tyr	Glu	Let
	Leu	Glu 130	Asn	Gly	Gln	Arg	Ala 135	Gly	Thr	Cys	Val	Leu 140	Glu	Tyr	Ala	Thr
35	Pro 145	Leu	Gln	Thr	Leu	Phe 150	Ala	Met	Ser	Gln	Туг 155	Ser	Gln	Ala	Gly	Phe 160
40	Ser	Gly	Glu	Asp	Arg 165	Leu	Glu	Gln	Ala	Lys 170	Leu	Phe	Cys	Arg	Thr 175	Leu
•	Glu	Asp	Ile	Leu 180	Ala	Asp	Ala	Pro	Glu 185	Ser	Gln	Asn	Asn	Cys 190	Arg	Leu
45	Ile	Ala	Тут 195	Gln	Glu	Pro	Ala	Asp 200	Asp	Ser	Ser	Phe	Ser 205	Leu	Ser	Gln
	Glu	Val 210	Leu	Arg	His	Leu	Arg 215	Gln	Glu	Glu	Lys	Glu 220	Glu	Val	Thr	Val
50	Gly 225	Ser	Leu	Lys	Thr	Ser 230	Ala	Val	Pro	Ser	Thr 235	Ser	Thr	Met	Ser	Gln 240
55	Glu	Pro	Glu	Leu	Leu 245	Ile	Ser	GÌy	Met	Glu 250	Lys	Pro	Leu	Pro	Leu 255	Arg
	Thr	Asp	Phe	Ser 260												

	(2)	INFO	DRMAT	NOI	FOR	SEQ	ID 1	<b>10</b> : 3	32:						*	
5 .			(i) : (xi)	() ()	A) L B) T D) T	ENGT YPE: OPOL	H: 4 ami: OGY:	8 am no a line	ino d cid ear	acid		: 33:	2 :			
10	Met 1	Thr	Pro	Gln	Lys 5	Pro	Ala	Leu	Ala	Val 10	Leu	Leu	Leu	Glu	Val 15	Pro
	Leu	Leu	Leu	Thr 20	Leu	Ser	Val	Leu	Lys 25	Lys	Arg	Cys	Leu	Va1 30	Thr	Cys
15	Glu	Pro	Thr 35	Ser	Arg	Phe	Val	Ser 40	Cys	Asp	Leu	Pro	Leu 45	Ser	Val	Xaa
20																
	(2)	INFO	ORMA	rion	FOR	SEQ	ID 1	NO: 3	33:							
25			(i) :	(.	A) L B) T	ENGT YPE:	H: 3 ami	ERIST 34 au no a lin	mino cid		ds					
30			(xi)					PTIO		EQ I	OM C	: 33	3:			
,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Met 1	Ala	Ala	Ala	Ala 5	Trp	Leu	Gln	Val	Leu 10	Pro	Val	Ile	Leu	Leu 15	Leu
35	Leu	Gly	Ala	His 20	Pro	Ser	Pro	Leu	Ser 25	Phe	Phe	Ser	Ala	Gly 30	Pro	Ala
	Thr	Val	Ala 35		Ala	Asp	Arg <sub>.</sub>	Ser 40	Lys	Trp	His	Ile	Pro 45	Ile	Pro	Ser
40	Gly	Lys 50	Asn	Tyr	Phe	Ser	Phe 55	Gly	Lys	Ile	Leu	Phe 60	Arg	Asn	Thr	Thr
45	Ile 65	Phe	Leu	Lys		Asp 70		Glu	Pro		Asp 75		Ser		Asn	
	Thr	Trp	Tyr	Leu	Lys 85	Ser	Ala	Asp	Cys	Туr 90	Asn	Glu	Ile	Tyr	Asn 95	Phe
50	Lys	Ala	Glu	Glu 100	Val	Glu	Leu	Tyr	Leu 105	Glu	Lys	Leu	Lys	Glu 110	Lys	Arg
	Gly	Leu	Ser 115	Gly	Lys	Tyr	Gln	Thr 120	Ser	Ser	Lys	Leu	Phe 125	Gln	Asn	Cys
55	Ser	Glu 130	Leu	Phe	Lys	Thr	Gln 135	Thr	Phe	Ser	Gly	Asp 140	Phe	Met	His	Arg
60	Leu 145	Pro	Leu	Leu	Gly	Glu 150	Lys	Gln	Glu	Ala	Lys 155	Glu	Asn	Gly	Thr	Asn 160

	Leu	Thr	Phe		Gly 165	Asp	Lys	Thr	Ala	Met 170	His	Glu	Pro	Leu	Gln 175	Thr
5	Trp	Gln	Asp	Ala 180	Pro	Tyr	Ile	Phe	Ile 185	Val	His	Ile	Gly	Ile 190	Ser	Ser
•	Ser	Lys	Glu 195	Ser	Ser	Lys	Glu	Asn 200	Ser	Leu	Ser	Asn	Leu 205	Phe	Thr	Met
10	Thr	Val 210	Glu	Val	Lys	Gly	Pro 215	Tyr	Glu	Tyr	Leu	Thr 220	Leu	Glu	Asp	Tyr
15	Pro 225	Leu	Met	Ile	Phe	Phe 230	Met	Val	Met	Cys	Ile 235	Val	Tyr	Val	Leu	Phe 240
	Gly	Val	Leu	Trp	Leu 245	Ala	Trp	Ser	Ala	Суs 250	Tyr	Trp	Arg	Asp	Leu 255	Leu
20	Arg	Ile	Gln	Phe 260		Ile	Gly	Ala	Val 265	Ile	Phe	Leu	Gly	Met 270	Leu	Glu
	_		275					280					285		Gly	
25		290	+				295					300				Lys
30	305	•		,		310					315	•				320
	Ile	· Val	Lys	Pro	325		Glu	Ser	Leu	Phe 330		Arg	Leu	ı Xaa		
35	(2)	INI	FORM	ATION	I FOE	R SEÇ	) ID	NO:	334:				,			
40	(2) INFORMATION FOR SEQ ID NO: 334:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 200 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 334:															
45		t Va 1	l Le	u Xaa		1 Va:	l Thi	c Le	ı Gly	/ Le		a Le	ı Ph	e Th	r Lei	ı Cys
	Gl	y Ly	s Ph	e Ly: 2		g Trj	) Ly:	s Le	u Ası 2!		y Ala	a Ph	e Le	u Le	u Ilo O	e Thr
50	Al	a Ph	e Le 3		r Va	l Le	u Il	e Tr		l Al	a Tr	p Me	t Th 4	r Me 5	t Ty	r Leu
55	Ph		y As O	n Va	l Ly	s Le	u Gl: 5		n Gl	y As	p Al	a Tr 6		n As	p Pr	o Thr
55		u Al	a Il	e Th	r Le	u Al 7		a Se	r Al	a Gl		r Se 5	r Se	r Se	r Se	r Thr 80
60	Pr	o Se	r Le	u Ar		r Th	r Al	a Pr	o Ph		s Gl	n Pr	о Су	s Ar	g Ar 9	g Thr 5

	Arg	Pro	Thr	Thr 100	Ser	Thr	Arg	Arg	Ser 105	Pro	Gly	Cys	Gly	Arg 110	Arg	Pro
5	Ser	Arg	Arg 115	Thr	Cys	Ser	Cys	Arg 120	Gly	Pro	Ile	Trp	Arg 125	Thr	Arg	Pro
10	Ser	Pro 130	Trp	Met	Asn	Thr	Met 135	Gln	Leu	Ser	Glu	Gln 140	Gln	Asp	Phe	Pro
	Thr 145	Ala	Ala	Trp	Glu	Lys 150	Asp	Pro	Val	Ala	Ala 155	Trp	Gly	Lys	Asp	Pro 160
15	Ala	Leu	Arg	Leu	Glu 165	Ala	Thr	Cys	Ile	Ser 170	Gln	Leu	Arg	Trp	Pro 175	Ser
	Cys	Ser	Thr	Val 180	Gly	Pro	Ser	Gln	Leu 185	Leu	Arg	Gln	Val	Thr 190	Gln	Glu
20	Xaa	Thr	Phe 195	Gly	Glu	Arg	Leu	Xaa 200								
					•											
25	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	10: 3	335:					=		
			(i)	(	A) L	ENGT	н: 2	4 am	ino		s					
30			(xi)	(	D) T	YPE: OPOL E DE:	OGY:	lin	ear	EQ II	D NO	: 33	5:			
35	Met 1	Leu	Leu	His	His 5	Gln	Leu	Leu	Ile	Val 10	Thr	Leu	His	Leu	Val 15	Leu
	Leu	Leu	Ala	Thr 20	Leu	Leu	Val	Xaa								
40	(2)	INF	ORMA'	rion	FOR	SEQ	ID 1	NO: :	336:							
se.			(i)	SEQU	ENCE	CHA	RACT	ERIS	rics	:						
45						engt Ype:				aci	ds					
			(xi)			OPOL E DE				EQ I	D NO	: 33	6:			
50	Met 1	Thr	Lys	Ala	Leu 5	Leu	Ile	Тут	Leu	Val 10	Ser	Ser	Phe	Leu	Ala 15	Leu
	Asn	Gln	Ala	Ser 20	Leu	Ile	Ser	Arg	Cys 25	Asp	Leu	· Ala	Gln	Val 30	Leu	Gln
55	Leu	Glu	Asp 35	Leu	Asp	Gly	Phe	Glu 40	Gly	Tyr	Ser	Leu	Ser 45	Asp	Trp	Leu
60	Cys	Leu 50		Phe	Val	Glu	Ser 55	Lys	Phe	Asn	Ile	Ser 60	Lys	Ile	Asn	Glu <sup>·</sup>

	Asn 65	Ala	Asp	Gly	Ser	Phe 70	qzA	Tyr	Gly	Leu	Phe 75	Gln	Ile	Asn	Ser	His 80
5	Tyr	Trp	Cys	Asn	Xaa 85	Tyr	Lys	Ser	Tyr	Ser 90	Glu	Asn	Leu	Cys	His 95	Val
	Asp	Cys	Gln	Asp 100	Leu	Leu	Asn	Pro	Asn 105	Leu	Leu	Ala	_	Ile 110	His	Cys
0	Ala	Lys	Arg 115	Ile	Val	Ser	Gly	Ala 120	Arg	Gly	Met	Asn	Asn 125	Trp	Val	Arg
15	Met	Glu 130	Xaa	Cys	Thr	Val	Gln 135	Ala	Gly	His	Ser	Ser 140	Thr	Gly	Xaa	
	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	10: 3	337:							
20			(i) :	(	A) L B) T	engt YPE :		5 am no a	ino cid	: acid	s			•		
25			(xi)	SEQ	UENC:	E DE	SCRI	PTIO	N: S	EQ I	ON O	: 33	7:			
	Met 1	Leu	Val	Ile	Ala 5	Gly	Gly	Ile	Leu	Ala 10	Ala	Leu	Leu	Leu	Leu 15	Ile
30	Val	Val	Val	Leu 20	Суѕ	Leu	Tyr	Phe	Lys 25	Ile	His	Asn	Ala	Leu 30	Lys	Ala
	Ala	Lys	Glu 35	Pro	Glu	Ala	Val	Ala 40	Val	Lys	Asn	His	Asn 45	Pro	Asp	Lys
35	Val	Trp 50	Trp	Ala	Lys	Asn	Ser 55	Gln	Ala	Lys	Thr	Ile 60	Ala	Thr	Glu	Ser
<b>1</b> 0	Cys 65	Pro	Ala	Leu	Gln	Cys 70	Cys	Glu	Gly	Tyr	Arg 75	Met	Суѕ	Ala	Ser	Phe 80
	Asp	Ser	Leu	Pro	Pro 85	Cys	Cys	Cys	Asp	Ile 90	Asn	Glu	Gly	Leu	Xaa 95	
<b>1</b> 5	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	vo: `:	338:		٠					
50				(	A) L B) T D) T	ENGT YPE : OPOL	H: 3 ami OGY:	8 am no a lin	cid ear	: acid EQ I		: 33	8:			
55	Met 1	Leu								-				Ala	Ala 15	Asn
	Val	Gly	Ala	Asn 20	Phe	Ala	Leu	Thr	Val 25	Glu	Lys	Ile	Gly	Met 30	Ile	Leu
50	Leu	Asn	Val	Ser	Glv	Хаа										

	$\cdot$	
5	(2) INFORMATION FOR SEQ ID NO: 339:	
10	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 39 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li><li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 339:</li></ul>	
15	Met Leu Val Val Ala Phe Gly Leu Leu Val Leu Tyr Ile Leu Leu Ala 1 5 10 15	
	Ser Ser Trp Lys Arg Pro Glu Pro Gly Ile Leu Thr Asp Arg Gln Pro 20 25 30	
20	Leu Leu His Asp Gly Glu Xaa 35	
25	(2) INFORMATION FOR SEQ ID NO: 340:	
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 71 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 340:	
	Ser Asp Pro Leu Ala Ser Ala Ser Gln Asn Ala Gly Ile Val Ser Val  1 5 10 15	
35	Gly Leu Cys Thr Arg Pro Gly Pro Gln Phe Lys Asn Ala Gln Pro Pro 20 25 30	
10	Phe Pro Xaa Gln Lys Ala Pro Arg Cys Leu Trp Glu Asn Gln Pro Pro 35 40 45	
	Pro Trp Arg Lys Ala Trp Asp Leu Pro Ser His Leu Gly Arg Arg Gly 50 55 60	
15	Ile Cys Gly Lys Ser Phe Xaa 65 70	
50	(2) INFORMATION FOR SEQ ID NO: 341:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 85 amino acids  (B) TYPE: amino acid	
55	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 341:	
	Tyr Val Met Ile Phe Lys Lys Glu Phe Ala Pro Ser Asp Glu Glu Leu 1 5 10 15	
50	Asp Ser The Are Are Clas Clas Clas Clas Dec Clas Law Ale Clas Clas	

•		٠		20					25					30		
_	Lys	Arg	Asn 35	Xaa	Lys	Glu	Leu	Ala 40	Gln	Arg	Gln	Xaa	Gly 45	Gly	Gly	Ser
5	Pro	Ala 50	Gly	Ala	Cys	Gly	Gly 55	Glu	Pro	Cys	Gln	Arg 60	Leu	Gln	Gly	Gln
10	Val 65	Gln	Pro	Pro	His	Arg 70	Gln	Gly	Ser	Ser	Gln 75	Arg	Arg	Ser	Pro	His 80
	Ala	Thr	Gly	Gln	Хаа 85		•									
15										÷						
	(2)	INFO	ORMA	rion	FOR	SEQ	ID I	NO: 3	342:							
20				(	A) L B) T D) T	ENGT YPE : OPOL	H: 9 ami OGY:	ERIST  O am  no a  lin  PTIO	ino cid ear	acid		: 34	2:			
25	Met 1	Trp	Asp	Trp	Asp 5	Trp	Ser	Ala	Pro	Trp 10	Ser	Trp	Pro	Leu	Trp 15	Leu
30	Ser	Leu	Ala	Leu 20	Val	Cys	Leu	Ser	Ala 25	Gly	Ala	Lys	Gly	His 30	Arg	Ala
	Ser	Glu	Ala 35	Gly	His	Ala	Arg	Ala 40	Leu	Thr	Cys	Glu	Met 45	Gly	Ser	Glu
35	Phe	Xaa 50	Thr	Ala	Xaa	Gly	<b>Leu</b> 55	Val	Leų	Gly	Xaa	<b>Xaa</b> 60	Xaa	Trp	Thr	Xaa
	Xaa 65	Asn	Gly	Ser	Ala	Gly 70	Pro	Glu	Arg	Arg	Gly 75	Trp	Arg	Pro	Ala	Ala 80
40	Phe	Leu	Ala	Val	Phe 85	Leu	Leu	Gly	Asp	Хаа 90						
45	(2)	INFO	RMAT	noi	FOR	SEQ	ID N	JO: 3	143:							
		1	(i) s	(2	A) L	ENGTI	H: 4	ERIST 8 am	ino a	acid	8					
50		•	(xi)	(1	TY (O	OPOL	OGY:	no ad lind PTION	ear		оио	: 34.	3:	,		-
55	Met 1	Phe	Gly	Pro	Thr 5	Phe	His	Ser	Leu	Val 10	Leu	Val	Pro	Pro	Trp 15	Pro
	Asn :	Leu	Ser	Leu 20	Leu	His	Phe	Thr	Ser 25	Pro	Val	Gly	Gln	His 30	Ser	Ser
60	Phe :	Leu	Pro 35	Thr	Ser	Leu	Arg	Leu 40	Xaa	Lys	Lys	Lys	Lys 45	Lys	Lys	Lys

5																
	(2)	INF	ORMAT	NOI	FOR	SEQ	ID i	vo: 3	344:							
10				(	A) L B) T D) T	CHAI ENGT YPE: OPOL E DE:	H: 5 ami OGY:	6 am no a lin	ino cid ear	acid		: 34	4:			
15	Met 1	Cys	Ser	Lys	Asn 5	Gly	Phe	Leu	Leu	Ala 10	Trp	Ser	Trp	Asn	Ser 15	Pro
20	Trp	Leu	Pro	Gln 20	Ala	Ser	Leu	Ala	His 25	Gly	Cys	Trp	Gly	Arg 30	Trp	Met
	Ser	Asp	Leu 35	Val	Gly	Cys	Ser	Arg 40	Glu	Asn	Lys	Cys	Ala 45	Leu	Arg	Asp
25	His	Ser 50	Glu	Arg	Val	Gln	Gly 55	Xaa			÷					
30	(2)			SEQUI	ENCE A) L	SEQ CHAI	RACTI H: 2	ERIS 22 a	rics mino		ds					
35		:	(xi)	(	D) T	YPE: OPOL E DE:	OGY:	lin	ear	EQ II	D NO	: 34	5:			
	Ser 1	Pro	Leu	Xaa	Phe 5	Cys	Val	Val	Leu	Leu 10	Leu	Gln	Ala	Ala	Arg 15	Gly
40	Tyr	Val	Val	Arg 20	Lys	Pro	Ala	Gln	Ser 25	Arg	Leu	Asp	Asp	Asp 30	Pro	Pro
45	Pro	Ser	Thr 35	. Leu	Leu	Lys	Asp	Tyr 40	Gln	Asn	Val	Pro	Gly 45	Ile	Glu	Lys
	Val	Asp 50	Asp	Val	Val	Lys	Arg 55	Leu	Leu	Ser	Leu	Glu 60	Met	Ala	Asn	Lys
50	Lys 65	Glu	Met	Leu	Lys	Ile 70	Lys	Gln	Glu	Gln	Phe 75	Met	Lys	Lys	Ile	Val 80
	Ala	Asn	Pro	Glu	Asp 85	Thr	Arg	Ser	Leu	Glu 90	Ala	Arg	Ile	Ile	Ala 95	Leu
55	Ser	Val	Lys	Ile 100	Arg	Ser	Tyr	Glu	Glu 105	His	Leu	Glu	Lys	His 110	Arg	Lys
						Arg										

	Lys	Met 130	Leu	Lys	Asn	Leu	Arg 135	Asn	Thr	Asn	Tyr	140	Val	Phe	Glu	Lys
5	Ile 145	Cys	Trp	Gly	Leu	Gly 150	Ile	Glu	Tyr	Thr	Phe 155	Pro	Pro	Leu	Tyr	Tyr 160
	Arg	Arg	Ala	His	Arg 165	Arg	Phe	Val	Thr	Lys 170	Lys	Ala	Leu	Суз	Ile 175	Arg
10	Val	Phe	Gln	Glu 180	Thr	Gln	Lys	Leu	Lys 185	Lys	Arg	Arg	Arg	Ala 190	Leu	Lys
15 <sup>.</sup>	Ala	Ala	Ala 195	Ala	Ala	Gln	Lys	Gln 200	Ala	Lys	Arg	Arg	Asn 205	Pro	Asp	Ser
	Pro	Ala 210	Lys	Ala	Ile	Pro	Lys 215	Thr	Leu	Lys	Asp	Ser 220	Gln	Xaa		
20 .	(2)	·	ORMA!	rion	FOR	SEQ	ID 1	NO: 3	346:							
25				(	A) L B) T D) T	ENGT YPE: OPOL	H: 6 ami OGY:	4 am no a lin	ino cid ear	acid		: 34	6:			
30	Met 1	Gly	Ala	Pro	Ala 5	Ala	Ser	Leu	Leu	Leu 10	Leu	Leu	Leu	Leu	Phe 15	Ala
	Cys	Cys	Trp	Ala 20	Pro	Gly	Ģly	, Ala	Asn 25	Leu	Ser	Gln	Asp	Asp 30	Ser	Gln
35	Pro	Trp	Thr 35	Ser	Asp	Glu	Thr	Val 40		Ala	Gly	Gly	Thr 45	Val	Val	Leu
40	Lys	Cys 50	GÌn	Val	Lys	Asp	His 55	Glu	Asp	Ser	Ser	Leu 60	Gln	Trp	Ser	Xaa
45	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: 3	347:							
50				(	A) L B) T D) T	ENGT YPE : OPOL	H: 1 ami OGY:	54 a no a lin	mino cid ear	aci		: 34	7:			. •
55	Met 1	Val	Ala	Pro	Val 5	Trp	Tyr	Leu	Val	Ala 10	Ala	Ala	Leu	Leu	Val 15	Gly
	Phe	Ile	Leu	Phe 20	Leu	Thr	Arg	Ser	Arg 25	Gly	Arg	Ala	Ala	Ser 30	Ala	Gly
60	<b>01</b>	<b>01</b>		_			<b>~</b> 3	<b>01</b>	•	.1-	<b>01</b>	<b>31</b> -	C1	3	**- 1	71-

			35					40					45			
5	Gln	Pro 50	Gly	Pro	Leu	Glu	Pro 55	Glu	Glu	Pro	Arg	Ala 60	Gly	Gly	Arg	Pro
J	Arg 65		Arg	Arg	Asp	Leu 70	Gly	Ser	Arg	Leu	Gln 75	Ala	Gln	Arg	Arg	Ala 80
10	Gln	Arg	Val	Ala	Trp 85	Ala	Glu	Ala	Asp	Glu 90	Asn	Glu	Glu	Glu	Ala 95	Val
	Ile	Leu	Ala	Gln 100	Glu		Glu	Gly	Val 105	Glu	Lys	Pro	Ala	Glu 110	Xaa	His
15	Leu	Ser	Gly 115	Lys	Ile	Gly	Ala	Lys 120	Lys	Leu	Arg	Xaa	Xaa 125	Glu	Glu	Lys
20	Gln	Ala 130	Arg	Lys	Ala	Gln	Xaa 135	Glu	Ala	Glu	Glu	Ala 140	Glu	Arg	Glu	Xaa
	Arg 145	Lys	Arg	Leu	Glu	Ser 150	Gln	Arg	Glu	Xaa						
25	(2)	INFO	ORMAT	rion	FOR	SEQ	ID N	ю: 3	148:							
			(i) !	SEOU	NCE.	CHAI	RACTI	7PT 91	יזרכ.	•						
30				() () ()	A) L: B) T D) T	ENGT YPE : OPOLA	H: 1' amin OGY:	7 am no a line	ino a cid ear	acid		: 348	3:			
	Met						Ser							Gln	Tm	Ser
35	1		_	_	5					10					15	
	Xaa														•	
40																
	(2)	INFO	RMAT	CION	FOR	SEQ	ID N	10: 3	49:							
45				() (I	A) LI 3) TY 0) TY	ENGTI YPE : OPOLO	RACTE H: 10 amir DGY: BCRIE	o ami no ac line	ino a cid ear	acids		349	):			
50	Met 1	Leu	Val	Cys	Ser 5	Phe	Leu	Phe	Leu	Xaa 10						
55	(2)	INFO	RMAT	lON	FOR	SEO	ID N	0: 3	50:							
							ACTE									

(A) LENGTH: 14 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 350:

```
Val Ile Glu Leu Cys Val Ser Leu Arg Ser Leu Asn Phe Xaa
 5
      (2) INFORMATION FOR SEQ ID NO: 351:
10
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 18 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 351:
15
      Met Cys Glu Phe Xaa Xaa Xaa Ile Met Xaa Leu Ala Gly Tyr Phe Ala
                        5
      Cys Xaa
20
      (2) INFORMATION FOR SEQ ID NO: 352:
25
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 62 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 352:
      Met Val Gly Gly Tyr Val Ser Ser Phe Ser Phe Pro Pro Val Ser Ser
                                           10
35
      Ser Leu Leu Pro Ala Ser Phe Ala Phe Pro Phe Leu Pro Gly Thr
      Pro Cys Pro Phe Leu Tyr Phe Leu Pro Ser Pro Phe Ser Pro Leu Pro
                                   40
40
      Leu Ser Leu Thr Arg Ser Asn Ser Phe Leu Leu Asn Gly Xaa
                               55
45
      (2) INFORMATION FOR SEQ ID NO: 353:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 33 amino acids
50
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 353:
      Glu Lys Lys Ser Met Ser Val Ser Asp Ile Tyr Ala Leu Glu Ser Leu
55
      Gly Arg Ser Leu Phe Thr Leu Asn Ser Met Cys Leu Pro Leu Ser Phe
                                       25
60
      Xaa
```

5	(2)	INF	CAMAC	rion	FOR	SEQ	ID N	10: 3	354:							
10			(i) :	- ( (	A) L B) T D) T	ENGT YPE : OPOL	H: 2 ami OGY:	45 a no a lin	mino cid ear	aci		: 35	<b>1</b> :			
15	Met 1	Gly	Gly	Ala	Ser 5	Arg	Arg	Val	Glu	Ser 10	Gly	Ala	Trp	Ala	Туг 15	Leu
13	Ser	Pro	Leu	Val 20	Leu	Arg	Lys		Leu 25	Glu	Ser	Leu	Val	Glu 30	Asn	Glu
20	Gly	Ser	Glu 35	Val	Leu	Ala	Leu	Pro 40	Glu	Leu	Pro	Ser	Ala 45	His	Pro	Ile
	Ile	Phe 50	Trp	Asn	Leu	Leu	Trp 55	Tyr	Phe	Gln	Arg	Leu 60	Arg	Leu	Pro	Ser
25	Ile 65	Leu	Pro	Gly	Leu	Val 70	Leu	Ala	Ser	Cys	Asp 75	Gly	Pro	Ser	Xaa	Ser 80
30	Gln	Ala	Pro	Ser	Pro 85	Trp	Leu	Thr	Pro	Asp 90	Pro	Ala	Ser	Val	Gln 95	Val
50	Arg	Leu	Leu	Trp 100	Asp	Val	Leu	Thr	Pro 105	Asp	Pro	Asn	Ser	Cys 110	Pro	Pro
35	Leu	Tyr	Val 115	Leu	Trp	Arg	Val	His 120	Ser	Gln	Ile	Pro	Gln 125	Arg	Val	Val
	Trp	Pro 130	Gly	Pro	Val	Pro	Ala 135	Ser	Leu	Ser	Leu	Ala 140	Leu	Leu	Glu	Ser
40	Val 145	Leu	Arg	His	Val	Gly 150	Leu	Asn	Glu	Val	His 155	Lys	Ala	Val	Gly	Leu 160
45	Leu	Leu	Glu	Thr	Leu 165	Gly	Pro	Pro	Pro	Thr 170	Gly	Leu	His	Leu	Gln 175	Arg
73	Gly	Ile	Tyr	Arg 180	Glu	Ile	Leu	Phe	Leu 185	Thr	Met	Ala	Ala	Leu 190	Gly	Lys
50	Asp	His	Val 195	Asp	Ile	Val	Ala	Phe 200	Asp	Lys	Lys	Tyr	Lys 205	Ser	Ala	Phe
	Asn	Lys 210	Leu	Ala	Ser	Ser	Met 215		Lys	Glu	Glu	Leu 220	Arg	His	Arg	Arg
55	Ala 225	Gln	Met	Pro	Thr	Pro 230	Lys	Ala	Ile	Asp	Cys 235	Arg	Lys	Cys	Phe	Gly 240
60	Ala	Pro	Pro	Glu	Cys 245											

	(2)	INF	ORMA'	TION	FOR	SEQ	ID	NO:	355:							
5			(i)	(	A) I B) 1		H: 3	5 an			ls	٠.				
10									N:S							
	Met 1	Lys	Phe	Ser	Leu 5		Phe	Leu	Pro	Met 10	Leu	Leu	Ile	Leu	Lys 15	Pro
15	Asp	Leu	Phe	His 20	Ile	Ser	Ile	Cys	Thr 25		Ala	Ala	Cys	Gly 30	Leu	Thr
	Phe	Pro	Хаа 35								.*					
20																
	(2)	INF	ORMA	rion	FOR	SEQ	ID i	NO:	356:							
25			(i)	(	A) I B) I		H: 2 ami	2 am no a			s	<i>\</i>				
			(xi)		-				N: S	EQ I	D NO	: 35	6:			
30	Met 1	Leu	Phe	Phe	Phe 5		Leu	His	Leu	Leu 10	Ser	Ile	Met	Ser	Phe 15	Leu
35	Ser	Pro	Asp	Ile 20	Met	Xaa				•					•	
55																
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: I	357:							
40				(	A) L B) T D) T	ENGT YPE: OPOL	H: 9 ami OGY:	8 am no a lin	ear	acid		25				
45			(xi)													
	Met 1	Phe	Gly	Leu	Leu 5	Val	Glu	Ser	Gln	Thr 10	Leu	Leu	Glu	Glu	Asn 15	Ala
50	Val	Gln	Gly	Thr 20	Glu	Arg	Thr	Leu	Gly 25	Leu	Asn	Ile	Ala	Pro 30	Phe	Ile
•	Asn	Gln	Phe 35	Gln	Val	Pro	Ile	Arg 40	Val	Phe	Leu	Asp	Leu 45	Ser	Ser	Leu
55	Pro	Cys 50	Ile	Pro	Leu	Ser	Lys 55	Pro	Val	Glu	Leu	Leu 60	Arg	Leu	Asp	Leu
60	Met 65	Thr	Pro	Tyr	Leu	Asn 70	Thr	Ser	Asn	Arg	Glu 75	Val	Lys	Val	Tyr	Val 80

PCT/US98/11422

548

Cys Xaa Ile Trp Glu Asp Leu Thr Ala Ile Pro Phe Trp Val Ser Tyr 90 Val Pro 5 (2) INFORMATION FOR SEQ ID NO: 358: 10 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 78 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 358: Met Phe Gly Ala His Arg Xaa Trp Gln Gly Ser Val Leu Leu Phe Leu 20 Ser Phe Ala Trp Gly Asn Gly Gly Ser Val Thr Phe Ser Asp Val Pro Arg Val Met Pro Leu Ala Gly Gly Pro Xaa Xaa Gln Val Ser Ser Thr 40 25 Pro Arg Pro Pro Pro His Gln Val Thr Ser Ser Pro Gly Leu Glu Ser Ala His Ile Val Cys Pro Glu Arg Lys Lys Lys Lys Lys 30 70 75 (2) INFORMATION FOR SEQ ID NO: 359: 35 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 359: Thr Leu Leu Xaa Phe Leu Xaa Leu Leu Thr Thr Glu Gly Gly Arg Glu 45 Asn Ile Phe Xaa Gly Arg Ile Leu Xaa Leu Gln Xaa Ser Pro Xaa 20 25 50 (2) INFORMATION FOR SEQ ID NO: 360: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 57 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 360: Met Leu Ser Phe Phe Ile Cys Leu Leu Ile Phe Val His Leu Leu Leu 5

(2) INFORMATION FOR SEQ ID NO: 364:

```
Leu Ser Phe Leu Ile Ser Asp Trp Pro Pro Pro Thr Gly Ser Ala Xaa
                  20 25
     His Lys Ile Leu Arg Leu Met Val Val Gln Arg Leu Ser Leu Leu Asp
 5
                                 40
     Gln Arg Lys Arg Trp Ser Glu Ala Xaa
10
      (2) INFORMATION FOR SEQ ID NO: 361:
             (i) SEQUENCE CHARACTERISTICS:
15
                    (A) LENGTH: 3 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 361:
20
     Lys Tyr Xaa
25
      (2) INFORMATION FOR SEO ID NO: 362:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 32 amino acids
                    (B) TYPE: amino acid
30
                   (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 362:
      Trp Ser Ser Ala Ser Ser Ser Trp Val Thr Thr Pro Glu Arg Ile Arg
                       5
35
      Pro Arg Met Asp Thr Leu Pro Val Lys Gly His Phe Leu Ser Met Xaa
                                    25
40
      (2) INFORMATION FOR SEQ ID NO: 363:
45
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 28 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
50
             (xi) SEQUENCE DESCRIPTION; SEQ ID NO: 363:
      Asp Ile Phe Val Phe Leu Leu Ser Thr Arg Ala Gly Gly Leu Gly Ile
                      5
55
      Asn Leu Thr Ala Xaa Asp Thr Val His Phe Leu Xaa
                  20
                                      25
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			(i)	SEQU	ENCE	: CHA	RACI	ERIS	TICS	:					•	
				(	(A) I	ENGI	H: 1	l5 an	nino	acid	ls					
_				(	(B) 3	YPE:	ami	ino a	cid							
5		,				OPOI										
			(xi)	SEQ	UENC	E DE	SCRI	PTIC	N: S	EQ I	D NC	): 36	4:			
	ml	*	<b>™</b>	0	D)	•	<b>03</b>	•		_		_		_		
	111	Leu	THE	Ser	Pne 5		GIU	Leu	Pro			Pro	GIU	Pro		
10	_				3					10					15	
	•															
	(2)	INF	ORMA	TION	FOR	SEO	ID :	NO:	365:							
	• •													,		
15			(i)	SEQU	ENCE	CHA	RACT	ERIS	TICS							
						ENGI					ls					
				(	B) 1	YPE:	ami	no a	cid							
				(	D) I	OPOL	OGY:	lin	ear							
••			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 36	5:			
20					•											
		His	Arg	Tyr	Ile	Thr	Phe	Phe	Lys	Cys	Phe	Arg	Ser	Val	Ile	Leu
	1				5					10					15	
	7.00	T 011	T 011	nh.	<b>71</b> -	T	G	D	<b>.</b>	~	~1		_	_,		_
25	rap.	Den	rea	20	116	Leu	Ser	PIO	25	ser	GIN	GIY	Cys		He	Leu
				2,0					23					30		
	Phe	Xaa														
30																
	(2)	INF	ORMA'	rion	FOR	SEQ	ID 1	NO:	366:							
			123	CEO! 1	- Con	C1173			<b></b>							•
35			(1)	SEQU.		CHA. ENGT					-					
, .						YPE:				acio	S					
						OPOL										
			(xi)	SEQ						EO I	D NO	: 36	6:			
				_						- <b>.</b>						
10	Met	Phe	Gly	Phe	Ile	Phe	Leu	Leu	Leu	Ile	Phe	Cys	Ile	Xaa	Leu	Cys
	1				5					10					15	
	_															
	Ser	Arg	Thr	Leu	Ser	Thr	Phe	Ile		Lys	Leu	Val	Gly	Phe	Leu	Tyr
15				20					25					30		
•	ш~~	T 2 00	Dho		<b>-</b> 1 -	<b>3</b>	T	O	•	•	<b>.</b>	-1	_			_
	ПĎ	Lys	35	Ser	TTE	ASII	Leu	ser 40	Leu	Leu	Leu	Thr	_	He	Lys	Lys
			,,,					40					45			
	Lys	Lvs	Lvs	Lys	Lvs	Lvs	Thr	Pro	Ara	Glv	Glv	Pro	Glv	Xaa	Gln	Ser
50	-	50			-,, -	-1-	55		9	<b>U</b> -3	<b>-</b> 1	60			· · · · ·	001
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	Pro	Pro														
	65															
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55																
	(2)		<b></b>													
	(2)	TNF	)KMA!	MOIT	FOR	SEQ	ID 1	VO: 3	367:							
			(3)	25017	-N/CVI-	CT I R	2 Y C-4771		DT	_						
50			(1)	SEQUI					rics		_					,

(B)	TYPE: amino acid	
(D)	TOPOLOGY: linear	

	•		(xi)	SEC	UENC	E DE	SCRI	PTIC	N: S	EQ 1	D NO	): 36	57:			
5	Met 1		Gly	' Leu	Gly 5		Pro	Arg	Gln	Ala 10		Trp	Thr	Leu	Met 15	Leu
10	Leu	Leu	Ser	Thr 20		Met	Tyr	Gly	Ala 25		Ala	Pro	Leu	Leu 30		. Leu
	Cys	His	Val 35		Gly	Arg	Val	Pro 40		Arg	Pro	Ser	Ser 45	Ala	Val	Leu
15	Leu	Thr 50	Glu	Leu	Thr	Lys	Leu 55		Leu	Cys	Ala	Phe 60	Ser	Leu	Leu	Val
	Gly 65		Gln	Ala	Trp	Pro 70	Gln	Gly	Pro	Pro	Pro 75	Trp	Arg	Gln	Ala	Ala 80
20	Pro	Phe	Ala	Leu	Ser 85	Ala	Leu	Leu	Tyr	Gly 90	Ala	Asn	Asn	Asn	Leu 95	Val
25	Ile	Tyr	Leu	Gln 100	Arg	Tyr	Met	Asp	Pro 105	Ser	Thr	Tyr	Gln	Val 110	Leu	Ser
			115					120			•		125			Arg
30		130					135					140				Ala
25	145				Tyr	150				•	155			•		160
35					Pro 165					170					175	
40				180	Gly				185					190		
			195		Val			200					205			
45		210			Gln		215					220				
	225				His	230					235					240
50	Gly	Phe	Ser	Gly	Trp \( 245	Ala	Ala	Leu	Val	Val 250	Leu	Ser	Gln	Ala	Leu 255	Asn
55	Gly	Leu	Leu	Met 260	Ser	Ala	Val	Met	Lys 265	His	Gly	Ser	Ser	Ile 270	Thr	Arg
		•	275		Ser			280					285			
60	Val	Leu 290	Leu	Arg	Leu	Gln	Leu 295	Thr	Ala	Ala	Phe	Phe 300	Leu	Ala	Thr	Leu

Leu Ile Gly Leu Ala Met Arg Leu Tyr Tyr Gly Ser Arg 5 (2) INFORMATION FOR SEQ ID NO: 368: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 31 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 368: 15 Met Gly Glu Gln Pro His Phe Ser Leu Cys Val Leu Leu Ala Ala Val 10 5 Arg Glu Asp Xaa Asp Pro Xaa Val Phe Pro Cys Cys Phe Leu Xaa 20 20 (2) INFORMATION FOR SEQ ID NO: 369: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 369: 30 Met Ser Phe Ile Ala Leu His Pro Leu Leu Pro Glu Ala Ala Leu Gly Val Pro Gly Gln Ser Pro His Arg Pro Leu Trp Gln Thr Gln Cys Cys 35 25 Val Ala Pro Pro Gln Pro Arg Ala Glu Phe Xaa 35 40 40 (2) INFORMATION FOR SEQ ID NO: 370: (i) SEQUENCE CHARACTERISTICS: 45 (A) LENGTH: 255 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 370: 50 Met Val Thr Ala Leu Thr Leu Leu Ala Phe Pro Leu Leu Leu His Ala Glu Arg Ile Ser Leu Val Phe Leu Leu Phe Leu Gln Ser Phe 25 55 Leu Leu Leu His Leu Leu Ala Ala Gly Ile Pro Val Thr Thr Pro Gly 40 Pro Phe Thr Val Pro Trp Gln Ala Val Ser Ala Trp Ala Leu Met Ala 60 50 55

•																
	Thr 65	Gln	Thx	Phe	Tyr	Ser 70	Thr	Gly	His	Gln	Pro 75	Val	Phe	Pro	Ala	Ile 80
5	His	Trp	His	Ala	Ala 85	Phe	Val	Gly	Phe	Pro 90	Glu	Gly	His	Gly	Ser 95	Cys
10	Thr	Trp	Leu	Pro 100	Ala	Leu	Leu	Val	Gly 105	Ala	Asn	Thr	Phe	Ala 110	Ser	His
	Leu	Leu	Phe 115	Ala	Val	Gly	Cys	Pro 120	Leu	Leu	Leu	Leu	Trp 125	Pro	Phe	Leu
15	Суѕ	Glu 130	Ser	Gln	Gly	Leu	Arg 135	Lys	Arg	Gln	Gln	Pro 140	Pro	Gly	Asn	Glu
	Ala 145	Asp	Ala	Arg	Val	Arg 150	Pro	Glu	Glu	Glu	Glu 155	Glu	Pro	Leu	Met	Glu 160
20	Met	Arg	Leu	Arg	Asp 165	Ala	Pro	Gln	His	Phe 170	Tyr	Ala	Ala	Leu	Leu 175	Gln
25	Leu	Gly	Leu	Lys 180	Tyr	Leu	Phe	Ile	Leu 185	Gly	Ile	Gln	Ile	Leu 190	Ala	Cys
	Ala	Leu	Ala 195	Ala	Ser	Ile	Leu	Arg 200	Arg	His	Leu	Met	Val 205	Trp	Lys	Val
30	Phe	Ala 210	Pro	Lys	Phe	Ile	Phe 215	Glu	Ala	Val	Gly	Phe 220	Ile	Val	Šer	Ser
	Val 225	Gly	Leu	Leu	Leu	Gly 230	Ile	Ala	Leu	Val	Met 235	Arg	Val	Asp	Gly	Ala 240
35	Val	Ser	Ser	Trp	Phe 245	Arg	Gln	Leu	Phe	Leu 250	Ala	Gln	Gln	Arg	Xaa 255	
40	(2)	INFO	RMAT	CION	FOR	SEQ	ID N	10: 3	371:							
			(i) s	(	A) L	CHAI ENGTI YPE:	H: 2	0 am	ino a		5					•
45			(xi)	(1	D) T	OPOLA E DES	OGY:	line	ear	EQ II	) NO:	: 371	l:			
50	Met 1	Xaa	Gly	Pro	Trp 5	Gly	Glu	Glu	Ala	Leu 10	Ile	Arg	Leu	Pro	Thr 15	Pro
	Ser	Gly	Leu	Хаа 20											,	•
55	(2)	INFO	RMAT	NOI	FOR	SEQ	ID N	ю: 3	72:							
		į	(i) S	SEQUE	ENCE	CHAI	RACTI	ERIST	rics:	: .						
60						engti Ype :				acids	5					

(B) TYPE: amino acid

```
(D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 372:
       Met Ala Thr Leu Glu Xaa Asn Gln Arg Glu Val Asp Arg Glu Ile Arg
  5
       Ser Leu Leu Trp Phe Leu Leu Cys Glu Ile Val Ser Gly Trp Leu
 10
       Cys Pro Glu Gly Pro Trp Phe Ser Gln Gly Cys Gln Ile Tyr Lys Asn
                                    40
       Leu Ser Ser Ser Ser Tyr Asn Leu Ser Phe Leu Leu Ser Leu Xaa
                                55
 15
20
       (2) INFORMATION FOR SEQ ID NO: 373:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 40 amino acids
25
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 373:
      Met Ile His Ser Gly Cys Thr Ser Gln Cys Leu Glu Gly Phe Phe Leu
30
                                           10
      Ile Phe Leu Leu Asp Phe Asn Pro Val Leu Ala Leu Asp Leu Ile Gly
                   20
35
      Ile Met Arg Lys Ala Ser His Xaa
               35
40
      (2) INFORMATION FOR SEQ ID NO: 374:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 35 amino acids
                    (B) TYPE: amino acid
45
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 374:
      Met Val Phe Ser Ala Arg Val Ser Leu Tyr Thr Arg Phe Lys Val Ile
50
     Leu Leu Ser Leu Leu Ile Met Ile Leu His Val Cys Trp Val Trp Val
      Ile Leu Xaa
55
              35
```

(2) INFORMATION FOR SEQ ID NO: 375:

```
(i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 11 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
  5
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 375:
       Gly Leu Leu Tyr Ile Met Tyr Cys Asn Ile Xaa
                        5
 10
       (2) INFORMATION FOR SEQ ID NO: 376:
              (i) SEQUENCE CHARACTERISTICS:
 15
                     (A) LENGTH: 64 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 376:
20
      Met Asn Asn Gly Leu Leu Gln Gln Pro Ser Ala Leu Met Leu Leu Pro
      Cys Arg Pro Val Leu Thr Ser Val Ala Leu Asn Ala Asn Phe Val Ser
                   20
                                       25
25
      Trp Lys Ser Arg Thr Lys Tyr Thr Ile Thr Pro Val Lys Met Arg Lys
      Ser Gly Gly Arg Asp His Thr Gly Gly Asn Lys Asp Arg Gly Ile Xaa
30
                               55
35
      (2) INFORMATION FOR SEQ ID NO: 377:
             (i) SEQUENCE CHARACTERISTICS:
40
                    (A) LENGTH: 19 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 377:
45
      Met Arg Lys Gln Arg Leu Val Pro Met Tyr Leu Gly Leu Ile Tyr Ile
        1
                                     10
      Leu Leu Xaa
50
      (2) INFORMATION FOR SEQ ID NO: 378:
55
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 5 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 378:
60
```

```
Met Arg Gln His Xaa
  5
       (2) INFORMATION FOR SEQ ID NO: 379:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 17 amino acids
 10
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 379:
       Leu Leu Pro Val Leu Ala Ser Ser Val Pro Ser His Ser Ala Thr
 15
                        5
                                           10
       Xaa
20
       (2) INFORMATION FOR SEQ ID NO: 380:
              (i) SEQUENCE CHARACTERISTICS:
25
                     (A) LENGTH: 84 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 380:
30
      Met Leu Pro Leu Leu Phe Thr Tyr Leu Asn Ser Phe Leu His Gln
      Arg Ile Pro Gln Ser Val Arg Ile Leu Gly Ser Leu Val Ala Ile Leu
                                       25
35
      Leu Val Phe Leu Ile Thr Ala Ile Leu Val Lys Val Gln Leu Asp Ala
      Leu Pro Phe Phe Val Ile Thr Met Ile Lys Ile Val Leu Ile Asn Ser
40
          50
      Phe Gly Ala Ile Leu Gln Gly Ser Leu Phe Gly Leu Ala Gly Leu Leu
                           70
45
      Pro Ala Ser Xaa
50 (2) INFORMATION FOR SEQ ID NO: 381:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 21 amino acids
                    (B) TYPE: amino acid
55
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 381:
      Met Lys Leu Ser Leu Phe Leu Ile Leu Ser Asp Val Phe Tyr Leu Gly
                        5
60
```

```
Ser Pro Xaa Thr Xaa
          20
```

5

- (2) INFORMATION FOR SEQ ID NO: 382:
  - (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 29 amino acids

10 (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 382:

Met Gly Thr Arg Arg Lys Gly Val Ala Trp Leu Ser Leu Ala Pro Leu 15 10

Ile Thr Gly Leu Ala Pro Ala His Ile Thr Ala Val Xaa 20 25

20

- (2) INFORMATION FOR SEQ ID NO: 383:
- (i) SEQUENCE CHARACTERISTICS: 25
  - (A) LENGTH: 34 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 383:
- 30 Met Lys Asp Leu Leu Gln Arg Asn Pro Trp Lys Asn Ser Leu Leu Leu 10

Leu Gln Val Cys Gln Ala Phe Leu Val Cys Ser Leu Thr Gln Leu Ala 20 25

35

Val Xaa

40

45

- (2) INFORMATION FOR SEQ ID NO: 384:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 47 amino acids
    - (B) TYPE: amino acid (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 384:
- Met Ser Glu Ser His Lys Ile Trp Trp Cys Tyr Arg His Leu Ala Phe 50 · 5 10

Pro Leu Leu Thr Leu Ile Leu Tyr Pro Ala Thr Leu Gly Arg Ser Val 25

- 55 Phe Cys His Asp Cys Lys Phe Pro Glu Ala Ser Pro Ala Met Xaa 40
- 60 (2) INFORMATION FOR SEQ ID NO: 385:

(2) INFORMATION FOR SEQ ID NO: 388:

```
(i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 25 amino acids
                      (B) TYPE: amino acid
  5
                      (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 385:
       Met Leu Asn Arg Ile Met Val Ala Ser Phe Gly Ala Val Leu Val Gln
 10
       Val Cys Arg Gly Xaa Gly Gln Gly Xaa
                    20
 15
       (2) INFORMATION FOR SEQ ID NO: 386:
              (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 68 amino acids
20
                      (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 386:
      Met Gln Leu Leu Leu Gly Leu Ile Arg Ser Gln Pro Ser Pro Pro
25
                                            10
      Pro Ser Leu Cys Leu Met Leu Cys Pro Cys Leu Pro Cys Leu Arg Tyr
                                       25
      Ser Pro Phe Val Pro Gln His Pro Cys Pro Leu Pro Leu Asp Leu Cys
30
                                   40
      Leu Ala Gly Cys Ser Ser Leu Ser Val Gln Asp Lys Cys Ser Trp Pro
                               55
35
      Tyr Pro Ile Xaa
       65
40
      (2) INFORMATION FOR SEQ ID NO: 387:
              (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 34 amino acids
45
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 387:
      Lys Glu Phe Phe Val Phe Leu Phe Val Cys Leu Phe Trp Leu Leu Ser
50
                        5
                                         10
      Asn Thr Pro Leu Thr Phe Ile Ser Ile Ile Leu Gln Arg Lys Glu Thr
                                      25
55
      Asn Xaa
```

		(i) SE	QUENCE C								
			(A) LEN								
<b>E</b>			(B) TYP								
5			QOT (Q)								
		(xi) S	EQUENCE !	DESCRI	PTION:	SEQ ID 1	NO: 3	88:			
	Ser Ph	e Leu M	et Val Le	eu Val	Ile Le	u Ala Al	la Se	r Pr	о Ха	a	
10	1		5			10					
10								•			
											•
	/2\ 7>*										
	(2) IN	FORMATI	ON FOR SE	Q ID N	Ю: 389	:					,
15		(i) an	^··								
13		(1) SE	QUENCE CH								
			(A) LENG								
			(B) TYPI								
		(rei) C	(D) TOPO	JLOGY:	linear						
20	•	(XI) S.	EQUENCE I	ESCRIP	PITON: S	SEQ ID N	io: 3	89:			
				•							
	1		5			10					
	-		,			10					
25											
	(2) IN	FORMATIC	N FOR SE	OIDNO	D: 390-						
		(i) SEQ	QUENCE CH	ARACTE	RISTICS	S:					
			(A) LENG								
30			(B) TYPE								
			(D) TOPO	LOGY:	linear						
		(xi) SE	QUENCE D	ESCRIP'	TION: S	EQ ID N	0: 39	0:			
35	Met Thr	Lys Al	a Arg Le	Phe A	Arg Leu	Trp Let	ı Val	Leu	Gly	Ser	Val
33	1		5			10				15	
	Dho Mar										
	rne met	i iie re	u Leu Ile	i Ile A	/al Tyr	Trp Asp	Ser	Ala	Gly	Ala	Ala
		2	0 ·		25				30		
40	His Dhe	Other to	n tria mba			_					
	mis rie	: <u>ryr</u> Le	u His Th	r Ser F	he Ser	Arg Pro	His		Gly	Pro	Pro
		33			40 .			45			
	Leu Pro	Thr Dr	o Clar Dec				_				
	50	*****	o Gly Pro	, MSD A	urg Asp	Arg GIU		Thr	Ala	Asp	Ser
45	30			55			60				
	Asp Val	Asp Xa	a Phe Lev	Acro V	'aa Dho	tau Cau				_	
	65	1-02-1-1	a Phe Leu 70		aa Pne			GIY	Val	Lys	
			, ,			75					80
	Ser Asp	Xaa Pro	Arg Lys	Glu T	hr Glu	Gln Pro	Dro	A1 =	D~o	C1	C
50	_		85	014 1	O.u	90	PLO	Ara	Pro		Ser
•						50				95	
	Met Glu	Glu Ser	Val Arg	Хаа Т	vr Asp	Tro Ser	Pro	Δτα	Yaa	λla	<b>7</b>
		100	)		105			AL 9	110	wia	Atg
					203				110		
55	Arg Thr	Gln Thi	Arg Ala	Gly S	er Xaa	Arg Xaa	Glv	Glv	Xaa	Cve	Cre
		115	_		20		1	125	-144	-73	~ys
	Gly Ala	Ser Ala	Pro Xaa	Pro A	la Trp	Pro Ser	Pro	Pro	Ara	Ser	Ala
<b>CO</b>	130			135	-		140				
60											

His Ser Thr Thr Ser Pro Thr Arg Ser Xaa 150 5 (2) INFORMATION FOR SEQ ID NO: 391: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 9 amino acids 10 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 391: Met Val Leu Leu Gly Leu Leu Ser Xaa . 15 5 (2) INFORMATION FOR SEQ ID NO: 392: 20 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 61 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 392: Met Cys Ile His Val Phe Met Xaa Val Leu Trp Val Leu Phe Leu Leu 30 Ash Pro Leu Cys Thr Gly Leu Trp Pro Leu Xaa Ash Cys Phe Ser Val Leu Arg His Ala Asp Trp Val Leu Gly Ala Asp Tyr Lys Gly Glu Glu 35 40 35 Leu Asn Arg His Gln Gly Pro Met Lys Pro Lys Asp Xaa 55. 60 40 (2) INFORMATION FOR SEQ ID NO: 393: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 447 amino acids 45 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 393: Met Leu Leu Gly Leu Leu Met Ala Ala Cys Phe Thr Phe Cys Leu Ser 50. His Gln Asn Leu Lys Glu Phe Ala Leu Thr Asn Pro Glu Lys Ser Ser 25 55 Thr Lys Glu Thr Glu Arg Lys Glu Thr Lys Ala Glu Glu Glu Leu Asp Ala Glu Val Leu Glu Val Phe His Pro Thr His Glu Trp Gln Ala Leu

	65 65	Pro	GIY	GIn	Ala	70	Pro	АТА	GIY	ser	75	vaı	Arg	ren	ast.	20
5	Gln	Thr	Gly	Glu	Arg 85	Glu	Ala	Lys	Leu	Gln 90	Tyr	Glu	qaA	Ŀys	Phe 95	Arg
	Asn	Asn	Leu	Lys 100	Gly	Lys	Arg	Leu	Asp 105	Ile	Asn	Thr	Asn	Thr 110	عات	Thr
10	Ser	Gln	Asp 115	Leu	Lys	Ser	Ala	Leu 120	Ala	Lys	Phe	Lys	Glu 125	Gly	Ala	Glu
15	Met	Glu 130	Ser	Ser	Lys	Glu	Asp 135	Lys	Ala	Arg	Gln	Ala 140	Glu	Val	Lys	Arg
	Leu 145	Phe	Arg	Pro	Ile	Glu 150	Glu	Leu	Lys	Lys	Asp 155	Phe	ÇZA	Glu	Leu	Asn 165
20	Val	Val	Ile	Glu	Thr 165	Asp	Met	Gln	Ile	Met 170	Val	Arg	Leu	Ile	Asn 175	≟ys
	Phe	Asn	Ser	Ser 180	Ser	Ser	Ser	Leu	Glu 185	Glu	Lys	Ile	Ala	Ala 190	Leu	Phe
25	Asp	Leu	Glu 195	Tyr	Tyr	Val	His	Gln 200	Met	Asp	Asn	Ala	Gln 205	Asp	Leu	Leu
30	Ser	Phe 210	Gly	Gly	Leu	Gln	Val 215	Val	Ile	Asn	Gly	Leu 220	Asn	Ser	<u> </u>	314
	Pro 225	Leu	Val	Lys	Glu	Tyr 230	Ala	Ala	Phe	Val	Leu 235	Gly	Ala	Ala	Phe	Ser 240
35	Ser	Asn	Pro	Lys	Val 245	Gln	Val	Glu	Ala	Ile 250	Glu	Gļy	Gly	Ala	Leu 255	3ln
	Lys	Leu	Leu	Val 260	Ile	Leu	Ala	Thr	Glu 265	Gln	Pro	Leu	Thr	Ala 270	Lys	Lys
40	Lys	Val	Leu 275	Phe	Ala	Leu	Cys	Ser 280	Leu	Leu	Arg	His	Phe 285	Pro	בעב	Ala
45	Gln	Arg 290	Gln	Phe	Leu	Lys	Leu 295	Gly	Gly	Leu	Gln	Val 300	Leu	Arg	Thr	Leu
	Val 305	Gln	Glu	Lys	Gly	Thr 310	Glu	Val	Leu	Ala	Val 315	Arg	Val	Val	- Time	ட் <b>e</b> ப 320
50	Leu	Tyr	Asp	Leu	Val 325	Thr	Glu	Lys	Met	Phe 330	Ala	Glu	Glu	Glu	Ala 335	Glu
	Leu	Thr	Gln	Glu 340	Met	Ser	Pro	Glu	Lys 345	Leu	Gln	Gln	Tyr	Arg 350	Glm	Val
55	His	Leu	Leu 355	Pro	Gly	Leu	Trp	Glu 360	Gln	Gly	Trp	Cys	Glu 365	Ile	Thr	Ala
60	His	Leu 370	Leu	Ala	Leu	Pro	Glu 375	His	Asp	Ala	Arg	Glu 380	Lys	Val	Lei	Gl'n

•	Thr Leu Gly Val Leu Leu Thr Thr Cys Arg Asp Arg Tyr Arg Gln Asp 385 390 395 400	
5	Pro Gln Leu Gly Arg Thr Leu Ala Ser Leu Glm Ala Glu Tyr Gln Val 405 410 415	
	Leu Ala Ser Leu Glu Leu Gln Asp Gly Glu Asp Glu Gly Tyr Phe Gln 420 425 430	
10	Glu Leu Leu Gly Ser Val Asn Ser Leu Leu Lys Glu Leu Arg Xaa 435 443 445	
15	(2) INFORMATION FOR SEQ ID NC: 394:  (i) SEQUENCE CHARACTERISTICS:	
20	(A) LENGTH: 24 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 394:	
25	Met Val Ile Ser Tyr Val Thr Phe Thr Pro Val Ser Ala Asp Cys Phe 1 5 10 15  Phe Asn Val Leu Val Cys Phe Kaa 20	
30	(2) INFORMATION FOR SEQ ID NC: 395:	
35	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 24 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 395:	
40	Glu Leu Leu Phe Leu Leu Ile Ile Ile Leu Gly Glu Ser Leu Ser Asp 1 5 10 15	
45	Val Ile Leu Ile Cys Phe <u>Kaa</u> 20	
,,,	(2) INFORMATION FOR SEQ ID NC: 396:	
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 35 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 396:	
55	Met Phe Tyr Trp Gly Gly Leu Ser Phe Tyr Phe Leu Leu Ser Ser Gly 1 5 10 15	
60	Val Gly Phe Tyr Cys Phe Leu Phe Gly Phe Gly Met Glu Ile Trp Ile 20 25 30	

```
Ala Ala Xaa
 5
      (2) INFORMATION FOR SEQ ID NO: 397:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 3 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 397:
     Gly Arg Xaa
15
     (2) INFORMATION FOR SEQ ID NO: 398:
20
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 25 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
25
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 398:
     Met Lys Leu Ser Leu Leu Ile Leu Thr Leu Met Gln Arg Tyr Phe Arg
                              10
30
     Thr Ile Thr Asn Ser Leu Cys Lys Xaa
                  20
35
     (2) INFORMATION FOR SEQ ID NO: 399:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 79 amino acids
                    (B) TYPE: amino acid
40
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 399:
     Met Pro Ala Val Ser Gly Pro Gly Pro Leu Phe Cys Leu Leu Leu Leu
45
     Leu Leu Asp Pro His Ser Pro Glu Thr Gly Cys Pro Pro Leu Arg Arg
     Phe Glu Tyr Lys Leu Ser Phe Lys Gly Pro Arg Leu Ala Leu Pro Gly
50
              35
                                  40
     Ala Gly Ile Pro Phe Trp Ser His His Gly Gly Glu Gly Gln Gly Trp
                              55
55
     Gly Pro Leu Cys Pro Gly Ser Leu Lys Val Leu Glu Gly Leu Xaa
```

60 (2) INFORMATION FOR SEQ ID NO: 400:

			(i) :													
					A) L					acid	s					
5					B) T D) T											
			(xi)							EO I	O NO	: 40	0:			
										-						
		Lys	Val	Phe	_	Ser	Met	Pro	Phe		Val	Leu	Phe	Gln		Leu
10	1				5					10					15	
10	τlρ	Gln	Glu	Acn	Yaa											
	-10	0	Olu	20	лыс								•			
1.5																
15	(2)	TATE	JOMAG	TON	EOD	CEO	TD 1	TO	101.							
	(2)	INP	ORMAT	LOW	FOR	SEQ	וטו	vo: 4	·UI:							
			(i) :	SEQUI	ENCE	CHA	RACT	ERIS'	rics	:						
				(.	A) L	ENGT	н: 2	57 a	mino	aci	ds .					
20					B) T											
			(xi)		D) די					FO T1	) NO	. 40	1.			
			(21)	SEQ.		. تارا د	JCI(I.		.v. 3	eQ I	) NO	. 40.	٠.			
	Met	Ala	Ala	Leu	Thr	Ser	His	Leu	Gln	Asn	Gln	Ser	Asn	Asn	Ser	Asn
25	1				5					10					15	
	Three or the	) co	Leu	7~~	Th~	7~~	Cor	Tiro	O.10	Tira	*	7.00	1707	Dha	Moh	Dua
	110	ASII	neu	20	TILL	Arg	261	цуз	25	цуs	Lys	ASD	vai	30	Mec	PLO
												•				
30	Pro	Ser	Ser	Ser	Ser	Glu	Leu		Glu	Ser	Arg	Gly		Ser	Asn	Phe
			35					40					45			
	Thr	Ser	Thr	His	Leu	Leu	Leu	Lvs	Glu	Aso	Glu	Glv	Val	Asp	Asp	Val
		50					55	-3-				60				
35																
	Asn 65	Phe	Arg	Lys	Val			Pro	Lys	Gly		Val	Thr	Ile	Leu	
	65					70	. •			-	75					80
	Gly	Ile	Pro	Ile	Lys	Lys	Thr	Lys	Lys	Gly	Cys	Arg	Lys	Ser	Cys	Ser
40					85				٠.	90					95	
		D1	••••	••	~	<b>.</b>		•	•	~1				_	_	
•	GIY	Pne	Val	100	Ser	Asp	Ser	гÀг	105	GIU	Ser	vai	Cys	Asn 110	Lys	Ala
						·			105					-10		
45	Asp	Ala	Glu	Ser	Glu	Pro	Val	Ala	Gln	Lys	Ser	Gln	Leu	Asp	Arg	Thr
			115					120					125			
	t/a1	C) re	Ile	50×	ð om	<b>71</b> ~	C111	21-	0.00	C1	C1		T ou	C0~	17-1	mb ~
	vai	130	116	Ser	ASD	Ala	135	AJA	Cys	GIY	Giu	140	Leu	ser	vai	1111
50														·		
		Glu	Glu	Asn	Ser	Leu	Val	Lys	Lys	Lys	Glu	Arg	Ser	Leu	Ser	Ser
	145					150					155					160
	ഭിഴ	Sor	Asn	Dhe	O.10	60×	Glu.	Cln	Laco	mb~	Co-	Gl.	Tla	T10	y can	Ties
55	~~y	JEI	uoii	- 11G	165	261	GIU	GIII	nys	170	Jer	GTĀ	***	***	175	₩,
	Phe	Cys	Ser		Lys	Asp	Ser	Glu		Asn	Glu	Lys	Tyr		Asp	Thr
				180					185					190		
60	Phe	Leu	Glu	Ser	Glu	Glu	Ile	Glv	Thr	Lvs	Val	Glu	Val	Val	Glu	Arg
-										_, _						5

			195					200					205			
5	Lys	Glu 210	His	Leu	His	Thr	Asp 215	Ile	Leu	Lys	Arg	Gly 220	Ser	Glu	Met	Asp
J	Asn 225	Asn	Cys	Ser	bio	Thr 230	Arg	Lys	Asp	Phe	Thir 235	Glu	Asp	Thr	Ile	Pro 240
10	Arg	Asn	Thr	Asp	Arg 245	Lys	Lys	Glu	Asn	Lys 250	Pro	Val	Phe	Phe	Gln 255	Gln
	Ile															
15																
20	(2)	INF		SEQUI	ENCE A) L B) T	CHA ENGT YPE:	ID N RACTI H: 4 ami: OGY:	ERIS 24 a no a	rICS mino cid		ds					
			(xi)				SCRI			EQ I	ON C	: <b>4</b> 0	2:			
25	Met 1	Glu	Lys	Gln	Cys 5	Суз	Ser	His	Pro	Val 10	Ile	Cys	Ser	Leu	Ser 15	Thr
30	Met	Tyr	Thr	Phe 20	Leu	Leu	Gly	Ala	Ile 25	Phe	Ile	Ala	Leu	Ser 30	Ser	Ser
	Arg	Ile	Leu 35	Leu	Val	Lys	Tyr	Ser 40	Ala	Asn	Glu	Glu	Asn 45	Lys	Tyr	Asp
35	Tyr	Leu 50	Pro	Thr	Thr	Val	Asn 55	Val	Суз	Ser	Glu	Leu 60	Val	Lys	Leu	Val
	Phe 65	Cys	Val	Leu	Val	Ser 70	Phe	Cys	Val	Ile	Lys 75	Lys	Asp	His	Gln	Ser 80
<b>40</b> .	Arg	Asn	Leu	Lys	Tyr 85	Ala	Ser	Trp	Lys	Glu 90	Phe	Ser	Asp	Phe	Met 95	Lys
45	Trp	Ser	Ile	Pro 100	Ala	Phe	Leu	Tyr	Phe 105	Leu	Asp	Asn	Leu	Ile 110	Val	Phe
	Tyr	Val	Leu 115	Ser	Tyr	Leu	Gln	Pro 120	Ala	Met	Ala	Val	Ile 125	Phe	Ser	Asn
50	Phe	Ser 130	Ile	Ile	Thr	Thr	Ala 135	Leu	Leu	Phe	Arg	Ile 140	Val	Leu	Lys	Xaa
	Arg 145	Leu	Asn	Trp	Ile	Gln 150	Trp	Ala	Ser	Leu	Leu 155	Thr	Leu	Phe	Leu	Ser 160
55	Ile	Val	Ala	Leu	Thr 165	Ala	Gly	Thr	Lys	Thr 170	Leu	Gln	His	Asn	Leu 175	Ala
60	Gly	Arg	Gly	Phe 180	His	His	Asp	Àla	Phe 185	Phe	Ser	Pro	Ser	Asn 190	Ser	Cys <sup>.</sup>

			195	3			-,-	200		-,0			205	••••	niu	цуз	
5	Glu	Trp 210	Thr	Phe	Pro	Glu	Ala 215	Lys	Trp	Asn	Thr	Thr 220	Ala	Arg	Val	Phe	
	Ser 225	His	Ile	Arg	Leu	Gly 230	Met	Gly	His	Val	Leu 235	Ile	Ile	Val	Gln	Cys 240	
10	Phe	Ile	Ser	Ser	Met 245	Ala	Asn	Ile	Tyr	Asn 250	Glu	Lys	Ile	Leu	Lys 255	Glu	
15	Gly	Asn	Gln	Leu 260	Thr	Glu	Xaa	Ile	Phe 265	Ile	Gln	Asn	Ser	Lys 270	Leu	Tyr	
	Phe	Phe	Gly 275	Ile	Leu	Phe	Asn	Gly 280	Leu	Thr	Leu	Gly	Leu 285	Gln	Arg	Ser	
20	Asn	Arg 290	Asp	Gln	Ile	Lys	Asn 295	Cys	Gly	Phe	Phe	Туг 300	Gly	His	Ser	Ala	
	Phe 305	Ser	Val	Àla	Leu	Ile 310	Phe	Val	Thr	Ala	Phe 315	Gln	Gly	Leu	Ser	Val 320	
25	Ala	Phe	Ile	Leu	Lys 325	Phe	Leu	Asp	Asn	Met 330	Phe	His	Val	Leu	Met 335	Ala	
30	Gln	Val	Thr	Thr 340	Val	Ile	Ile	Thr	Thr 345	Val	Ser	Val	Leu	Val 350	Phe	Asp	
	Phe	Arg	Pro 355	Ser	Leu	Glu	Phe	Phe 360	Leu	Glu	Ala	Pro	Ser 365	Val	Leu	Leu	
35	Ser	Ile 370	Phe	Ile	Tyr	Asr.	Ala 375	Ser	Lys	Pro	Gln	Val 380	Pro	Glu	Tyr	Ala	
	Pro 385	Arg	Gln	Glu	Arg	Ile 390	Arg	Asp	Leu	Ser	Gly 395	Asn	Leu	Trp	Glu	Arg 400	
<b>1</b> 0	Ser	Ser	Gly	Asp	Gly 405	Glu	Glu	Leu	Glu	Arg 410	Leu	Thr	Lys	Pro	Lys 415	Ser	
15	Asp	Glu	Ser	Asp 420	Glu	Asp	Thr	Phe									
	(2)	INFO	ORMAT	NOI	FOR	SEQ	ID N	IO: 4	103 :								
50		•	(i) :	(1	A) LI B) T	ENGT YPE :	H: 3	3 am	ino a cid		s						
55			(xi)	SEQ		OPOLA E DES				EQ II	ON C	: 403	3:				
	Met 1	Trp	Gly	Gln	Gly 5	Ser	Gln	Lys	Ser	His 10	Phe	Ser	Asp	Leu	Val 15	Phe	
50	Gly	Val	Arg	Glu 20	Leu	Cys	Ala	Gln	Pro 25	Ser	Asp	Pro	Gly	Ser 30	Pro	His	

PCT/US98/11422

Xaa

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12	INFORMATION	EVOD	CEO	TD	MO.	101
\ Z	) INCOMPLION	FUR	SEU	עג	MO:	404

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 80 amino acids

- (B) TYPE: amino acid
- (D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 404:
- 15 Met Val Gln His Ile Gln Pro Ala Ala Leu Ser Leu Leu Ala Gln Trp
  1 5 10 15
  - Ser Thr Leu Val Gln Glu Leu Glu Ala Ala Leu Gln Leu Ala Phe Tyr 20 25 30
  - Pro Asp Ala Val Glu Glu Trp Leu Glu Glu Asn Val His Pro Ser Leu 35 40 45
- Gln Arg Leu Gln Xaa Leu Leu Gln Asp Leu Ser Glu Val Ser Ala Pro 55 60
  - Pro Leu Pro Pro Thr Ser Pro Gly Arg Asp Val Ala Gln Asp Pro Xaa 65 70 75 80

30

- 35 (2) INFORMATION FOR SEQ ID NO: 405:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 95 amino acids
    - (B) TYPE: amino acid
- 40 (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 405:

Met Leu Asn Gln Gly Tyr Ile Arg Lys Ile Ile Leu Ile Ile Ile Leu

1 5 10 15
45

Gly Ser Phe Ser Ser Pro Lys Lys Ala Ile Leu Met Gly Phe Gln Asn 20 25 30

Gln Lys Lys Ala Leu Asn Glu Glu Gln Thr Thr Gly Val Pro Met Ser 50 35 40 45

Ile Ser Gly Lys Leu Arg Pro Ser Arg Ser Leu Asp Phe Val Gln Pro 50 55 60

Pro Arg Phe Gln Ser Gln Gln Pro Ser Ala Val Val Asp Arg Arg Gly 65 70 75 80

Phe Xaa Xaa Lys Ala Ala Arg Gly Gln Glu Phe Ser Glu Ser Xaa 85 90 95

PCT/US98/11422

	(2)	INF	ORMAT	NOI	FOR	SEQ	ID 1	<b>10</b> : 4	106:							
			(i) :	(	A) L B) T	CHAI ENGT: YPE:	H: 2 ami	57 a no a	mino cid		ds		•			
10			(xi)							EQ II	ON O	: 40	6:			
10	Met 1	_	Gly	Pro	Ala 5	Gln	Ala	Lys	Leu	Leu 10	Pro	Gly	Ser	Ala	Ile 15	Gln
15	Ala	Leu	Val	Gly 20	Leu	Ala	Arg	Pro	Leu 25	Val	Leu	Ala	Leu	Leu 30	Leu	Val
	Ser	Ala	Ala 35	Leu	Ser	Ser	Val	Val 40	Ser	Arg	Thr	Asp	Ser 45	Pro	Ser	Pro
20	Thr	Val 50	Leu	Asn	Ser	His	Ile 55	Ser	Thr	Pro	Asn	Val 60	Asn	Ala	Leu	Thr
25	His 65	Glu	Asn	Gln	Thr	Lys 70	Pro	Ser	Ile	Ser	Gln 75	Ile	Ser	Thr	Thr	Leu 80
	Pro	Pro	Thr	Thr	Ser 85	Thr	Lys	Lys	Ser	Gly 90	Gly	Ala	Ser	Val	Val 95	Pro
30	His	Pro	Ser	Pro 100	Thr	Pro	Leu	Ser	Gln 105	Glu	Glu	Ala	Asp	Asn 110	Asn	Glu
	Asp	Pro	Ser 115	Ile	Glu	Glu	Glu	Asp 120	Leu	Leu	Met	Leu	Asn 125	Ser	Ser	Pro
35	Ser	Thr 130	Ala	Lys	Asp	Thr	Leu 135	Asp	Asn	Gly	Asp	Tyr 140	Gly	Glu	Pro	Asp
40	Tyr 145	Asp	Trp	Thr	Thr	Gly 150	Pro	Arg	qaA	Asp	Asp 155	Glu	Ser	Asp	Asp	Thr 160
70	Leu	Glu	Glu	Asn	Arg 165	Gly	Тут	Met	Glu	Ile 170	Glu	Gln	Ser	Val	Lys 175	Ser
45	Phe	Lys	Met	Pro 180	Ser	Ser	Asn	Ile	Glu 185	Glu	Glu	Asp	Ser	His 190	Phe	Phe
	Phe	His	Leu 195	Ile	Ile	Phe	Ala	Phe 200	Суѕ	Ile	Ala	Val	Val 205	Tyr	Ile	Thr
50	Tyr	His 210	Asn	Lys	Arg	Lys	Ile 215	Phe	Leu	Leu	Val	Gln 220	Ser	Arg	Lys	Trp
55	Arg 225	Asp	Gly	Leu	Cys	Ser 230	Lys	Thr	Val	Glu	Tyr 235	His	Arg	Leu	Asp	Gln 240
33	Asn	Val	Asn	Glu	Ala 245	Met	Pro	Ser	Leu	Lys 250	Ile	Thr	Asn	Asp	Tyr 255	Ile

Phe

5	(2)	INFO	ORMAT	NOI	FOR	SEQ	ID N	10: 4	107 :							
,			(i) :	(	A) L B) T	ENGT YPE:	H: 6	23 a no a	mino cid		ds					
10			(xi)	SEQU			OGY: SCRI			EQ II	оис	: 40	7:			
	Met 1	Phe	Met	Arg	Ile 5	Ala	Lys	Ala	Tyr	Ala 10	Ala	Leu	Thr	Asp	Glu 15	Ğlu
15	Ser	Arg	Lys	Asn 20	Trp	Glu	Glu	Phe	Gly 25	Asn	Pro	Asp	Gly	Pro 30	Gln	Ala
20	Thr	Ser	Phe 35	Gly	Ile	Ala	Leu	Pro 40	Ala	Trp	Ile	Val	Asp 45	Gln	Lys	Asn
	Ser	Ile 50	Leu	Val	Leu	Leu	Val 55	Tyr	Gly	Leu	Ala	Phe 60	Met	Val	Ile	Leu
25	Pro 65	Val	Val	Val	Gly	Ser 70	Trp	Trp	Tyr	Arg	Ser 75	Ile	Arg	Tyr	Ser	80
	Asp	Gln	Ile	Leu	Ile 85	Arg	Thr	Thr	Gln	Ile 90	Tyr	Thr	Tyr	Phe	Val 95	Tyr
30	Lys	Thr	Arg	Asn 100	Met	Asp	Met	Lys	Arg 105	Leu	Ile	Met	Val	Leu 110	Xaa	Gly
35	Ala	Ser	Glu 115	Phe	Asp	Pro	Gln	Тут 120	Asn	Lys	Asp	Ala	Thr 125	Ser	Arg	Pro
	Thr	Asp 130	Asn	Ile	Leu	Ile	Pro 135		Leu	Ile	Arg	Glu 140	Ile	Gly	Ser	Ile
40	Asn 145	Leu	Lys	Lys	Asn	Glu 150	Pro	Pro	Leu	Thr	Cys 155	Pro	Tyr	Ser	Leu	Lys 160
	Ala	Arg	Val	Leu	Leu 165	Leu	Ser	His	Leu	Ala 170	Arg	Met	Lys	Ile	Pro 175	Glu
45	Thr	Leu	Glu	Glu 180	Asp	Gln	Gln	Phe	Met 185	Leu	Lys	ГЛЗ	Cys	Pro 190	Ala	Leu
50	Leu	Gln	Glu 195	Met	Val	Asn	Val	Ile 200	Cys	Gln	Leu	Ile	Val 205	Met	Ala	Arg
	Asn	Arg 210		Glu	Arg	Glu	Phe 215	Arg	Ala	Pro	Thr	Leu 220	Ala	Ser	Leu	Glu
55	Asn 225	Ċys	Met	Lys	Leu	Ser 230	Gln	Met	Ala	Val	Gln 235	Gly	Leu	Gln	Gln	Phe 240
	Lys	Ser	Pro	Leu	Leu 245	Gln	Leu	Pro	His	Ile 250	Glu	Glu	Asp	Asn	Leu 255	Arg
60	Arg	Val	Ser	Asn	His	Lys	Lys	Tyr	Lys	Ile	Lys	Thr	Ile	Gln	Asp	Leu

•				260					265					270		
5	Val	Ser	Leu 275	Lys	Glu	Ser	Asp	Arg 280	His	Thr	Leu	Leu	His 285	Phe	Leu	Gli
	Asp	Glu 290	Lys	Tyr	Glu	Glu	Val 295	Met	Ala	Val	Leu	Gly 300	Ser	Phe	Pro	Ту
10	Val 305	Thr	Met	Asp	Ile	Lys 310	Ser	Gln	Val	Leu	Asp 315	Asp	Glu	Asp	Ser	As: 320
	Asn	Ile	Thr	Val	Gly 325	Ser	Leu	Val	Thr	Val 330	Leu	Val	Lys	Leu	Thr 335	Arg
15	Gln	Thr	Met	Ala 340	Glu	Val	Phe	Glu	Lys 345	Glu	Gln	Ser	Ile	Cys 350	Ala	Ala
20	Glu	Glu	Gln 355	Pro	Ala	Glu	Asp	Gly 360	Gln	Gly	Glu	Thr	Asn 365	Lys	Asn	Arg
	Thr	Lys 370	Gly	ĠĺŊ	Trp	Gln	Gln 375	Lys	Ser	Lys	Gly	Pro 380	Lys	Lys	Thr	Ala
25	Lys 385	Ser	Lys	Lys	Lys	Lys 390	Pro	Leu	Lys	Lys	Lys 395	Pro	Thr	Pro	Val	Le:
	Leu	Pro	Gln	Ser	Lys 405	Gln	Gln	Lys	Gln	Lys 410	Gln	Ala	Asn	Gly	Val 415	Va.
30	Gly	Asn	Glu	Ala 420	Ala	Val	Lys	Glu	Asp 425	Glu	Glu	Glu	Val	Ser 430	Asp	Lys
35	Gly	Ser	Asp 435	Ser	Glu	Glu	Glu	Glu 440	Thr	Asn	Arg	Asp	Ser 445	Gln	Ser	Glu
		450		Gly			455.					460		_		_
<b>4</b> 0	Gln 465	Asn	Lys	Asp	Asp	Glu 470	Ala	Glu	Trp	Gln	Glu 475	Leu	Gln	Gln	Ser	11e
4.5				Glu	485				•	490			_		495	
<b>1</b> 5				Ser 500					505					510		
50	Leu	Tyr	Ile 515	Ala	Asp	Arg	Lys	Glu 520	Gln	Thr	Leu	Ile	Ser 525	Met	Pro	Туз
		530		Thr		_	535					540		-		
55	Ala 545	Pro	Gly	Lys	Pro	Gly 550	Asn	Tyr	Gln	Tyr	Thr 555	Val	Phe	Leu	Arg	Se:
<b>.</b>	Asp	Ser	Tyr	Met	Gly 565	Leu	qeA	Gln	Ile	Lys 570	Pro	Leu	Glu	Val	Xaa 575	Lys
50	Phe	Met	Arg	Leu	Lys	Pro	Val	Pro	Glu	Asn	His	Pro	Gln	Trp	Asp	Thi

•				580					585					590		
5	Ala	Ile	Glu 595	Gly	Asp	Glu	Asp	Gln 600	Glu	Asp	Ser	Glu	Gly 605	Phe	Glu	Asp
	Ser	Phe 610	Glu	Gly	Gly	Arg	Gly 615	Arg	Glu	Glu	Gly	Arg 620	Trp	Trp	Thr	
10	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	vo: 4	408:		٠					
15				(	A) L B) T D) T	CHAI ENGT YPE: OPOL E DE:	H: 1 ami OGY:	90 a no a lin	mino cid ear	aci		: 40	8:			
20	Met 1	Lys	Ala	Ser	Gln 5	Cys	Cys	Cys	Cys	Leu 10	Ser	His	Leu	Leu	Ala 15	Ser
	Val	Leu	Leu	Leu 20	Leu	Leu	Leu	Pro	Glu 25	Leu	Ser	Gly	Xaa	Leu 30	Xaa	Val
25	Leu	Leu	Gln 35	Ala	Ala	Glu	Ala	Ala 40	Pro	Gly	Leu	Gly	Pro 45	Pro	Asp	Pro
30	Arg	Pro 50	Arg	Thr	Leu	Pro	Pro 55	Leu	Pro	Pro	Gly	Pro 60	Thr	Pro	Ala	Gln
	Gln 65	Pro	Gly	Arg	Gly	Leu 70	Ala	Glu	Ala	Ala	Gly 75	Pro	Arg	Gly	Ser	Glu 80
35	Gly	Gly	Asn	Gly	Ser 85	Asn	Pro	Val	Ala	Gly 90	Leu	Glu	Thr	Asp	Asp 95	His
	Gly	Gly	Lys	Ala 100	Gly	Glu	Gly	Ser	Val 105	Gly	Gly	Gly	Leu	Ala 110	Val	Ser
40	Pro	Asn	Pro 115	Gly	Asp	Lys	Pro	Met 120	Thr	Gln	Arg	Ala	Leu 125	Thr	Val	Leu
45	Met	Val 130	Val	Ser	Gly	Ala	Val 135	Leu	Val	Tyr	Phe	Val 140	Val	Arg	Thr	Val
	Arg 145	Met	Arg	Arg	Arg	Asn 150	Arg	Lys	Thr	Arg	Arg 155	Tyr	Gly	Val	Leu	Asp 160
50	Thr	Asn	Ile	Glu	Asn 165	Met	Glu	Leu	Thr	Pro 170	Leu	Glu	Gln	Asp	Asp 175	Glu
	Asp	Asp	Asp	Asn 180	Thr	Leu	Phe	Asp	Ala 185	Asn	His	Pro	Arg	Arg 190		
55				•					-							
	(2)	INF	ORMA!	ron	FOR	SEQ	ID I	10: 4	<b>1</b> 09 :							

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 179 amino acids

•						yfe: Cycl										
			(xi)			E DE				EQ II	011	: 40	9:			
5	Met 1	Ser	Pro	Ser	G1;·	Arg	Leu	Cys	Leu	Leu 10	Thr	Ile	Val	Gly	Leu 15	Il
0	Leu	Pro	Thr	20 20	Gly	Gln	Thr	Leu	1:/s 25	Asp	Thr	Thr	Ser	Ser 30	Ser	Se
	Ala	ązĄ	Ser 35	īhr	Ile	Met	Asp	Ile 40	Gln	Val	Pro	Thr	Arg 45	Ala	Pro	Ası
15	Ala	Val 50	īyr	Thr	Glu	Leu	Gln 55	₽±o	Thr	Ser	Pro	Thr 60	Pro	Thr	<u>arb</u>	Pr
	Ala 65	qzA	Glu	Ipr	Pro	GLn 70	Pro	Gjip	The	Gln	Thr 75	Gln	Gln	Leu	Glu	Gl; 8
20	Thr	qzA	Gly	Sto	Leu 85	Val	The	Asp	Pro	Glu 90	Thr	His	Lys	Ser	Thr 95	Ly
25	Ala	Ala	His	Pro 100	Thr	೫ವರಿ	ÇZA	Thr	Thr 105	Thr	Leu	Ser	Glu	Arg 110	Pro	Se
	Pro	Ser	Thr 115	Ąsp	Val	GLn	Thr	А <del>з</del> р 120	Pro	Gln	Thr	Leu	Lys 125	Pro	Ser	Gl:
30	Phe	His 130	Glu	Yeż	Asp	3≈0	Phe 135	Pie	Zva	Asp	Glu	His 140	Thr	Leu	Arg	Ly
_	Arg 145	Gly	Leu	Leu	Val	Ala 150	Ala	Val	Leu	Phe	Ile 155	Thr	Gly	Ile	Ile	11
35	Leu	Thr	Ser	Gly	Lys 165	C::s	Āŗg	Gln	Leu	Ser 170	Arg	Leu	Cys	Arg	Asn 175	Hi
10	Cys	Arg	Kaa													
	(2)	INFO	CRMA:	PICN	PĊF.	SEQ	ID I	NC: 4	110:							
15			(i) .	· (	A) 1 3) 1	CEA Engl IFE: CFOL	H: 1 ami	4 am no a	ino cid		s					
50	Met					E DE Gln				_				Xaa		
55	1		_,	-,-3	5					10						
, ,	(2)	INF	ORMA!	TION	FCR	SEQ	ID I	.vo: 4	111:							

(i) SEQUENCE CHARACTERISTICS:

60

(A) LENGTH: 232 amino acids

(B) TYPE: amino acid

			(xi)	) SEQ			LOGY ESCRI			SEQ 1	D NO	D: 41	11:			
5	Met	: Lev	ı Ala	Gly	Lys 5		ılle	Pro	Val	. His		Val	. Arg	Gly	Leu 15	Lys
	Glu	ı Lys	; Ile	Val 20	Arg	Ser	Phe	: Glu	Val 25		Pro	Asp	Gly	Ser 30		e Leu
10	Lev	ı Ile	Asn 35	Gly	Ile	Ala	Gly	Тут 40		His	Leu	Leu	Ala 45	Met	: Lys	Thr
15.	Lys	50 50	Leu	Ile	Gly	Ser	Met 55		Ile	Asn	Gly	Arg 60	Val	Ala	Ala	Ser
	65					70					75					Gly 80
20					85					90					95	Phe
				100					105					110		Asn
25			115					120					125			Tyr
30		130					135					140				Ala
	145		Asn			150					155					160
35			Ile		165					170					175	
40			His	180					185					190		
40			Lys 195					200					205		•	
45		210	Tyr				215		Glu	Lys	Gly	Lys 220	Ala	Leu	Met	Tyr
	225	ren	His	His		Ser 230	Asp	Phe								
50	(2)	INFO	RMAT	'ION	FOR	SEQ	ID N	o: 4	12:							
55			(i) s	(E (E	L) LE 3) TY 3) TO	NGTI PE: POLO	f: 54 amin XGY:	l ami o ac line	no a id ar	cids		412	:			•
60	Ile 1	Leu	Leu :	Cys :	Ser :	l'rp	Pro '	Thr	Gly :	Leu ' 10	Val (	Gly (	Gly a	Arg .	Asp 15	Pro

	Gly	Ser	Ser	Arg 20	Gly	Ser	Ser	Ala	Ser 25	Leu	Thr	Pro	Ser	Pro.	Ġly	Arg
5	Gln	Pro	Cys 35		Arg	Arg	Arg	Gly 40	Tyr	Ser	Val	Gly	Arg 45	Arg	Ser	Ser
10	Pro	Pro 50	Asp	Gly	Ser	Xaa										
	(2)	INFO	ORMAT	rion	FOR	SEO	ID I	NO: 4	413:							
15			(i) :	SEQUI () ()	ENCE A) L B) T D) T	CHAI ENGT YPE: OPOL	RACT H: 3 ami OGY:	ERIST 3 am no a lin PTIO	TICS ino cid ear	acid		: 41:	3:			
20	Met 1	Ser	Leu	Gln	Ser 5	Asn	Ala	Trp	Ser	Lys 10	Xaa	Leu	Phe	Ile	Val 15	Phe
25	Leu	Phe	Leu	Arg 20	Val	Leu	Phe	Lys	Thr 25	Gly	Val	Ser	Ser	Glu 30	Glu	Ser
	Xaa				•											
30																
	(2)	TNIC	ר אינו	ntost	EOD	CEO.	TD 1	v.	414.							
35	(2)		(i) :	SEQUI () ()	ENCE A) L B) T D) T	CHAI ENGT YPE: OPOL	RACT H: 2 ami OGY:	NO: 4 ERIS 19 a no a lin PTIO	TICS mino cid ear	aci		: 41	<b>4</b> :			
35 40	•		(i) : (xi)	SEQUI () () () SEQU	ENCE A) L B') T D) T UENC	CHAI ENGT YPE: OPOL E DE	RACT H: 2 ami OGY: SCRI	ERIS' 19 a no a lin	TICS mino cid ear N: S	aci EQ I	D NO			Ser	Leu 15	Ala
	Met 1 Ala	Ala Arg	(i) (xi) Val Ala	SEQUI ( ( SEQUI Val Ile 20	ENCE A) L B) T D) T UENC: Leu 5	CHAI ENGT YPE: OPOL E DE: Leu Val	RACT H: 2 ami OGY: SCRI Ala	ERIS' 19 a no a lin PTIO Asn	TICS mino cid ear N: S Leu Gly 25	aci EQ I Ala 10 Ser	D NO Gln Ile	Gly Gly	Asp Asn	Leu 30	15 Leu	Gly
40	Met 1 Ala	Ala Arg	(i) : (xi) Val Ala Glu 35	SEQUI () () SEQUI Val Ile 20	ENCE A) L B) T D) T UENC: Leu 5 Ala Ser	CHAI ENGT YPE: OPOL E DE Leu Val	RACT H: 2 ami OGY: SCRI Ala Gln	ERIS' 19 a no a lin PTIO Asn Lys Ala 40	TICS mino cid ear N: S Leu Gly 25 Thr	aci EQ II Ala 10 Ser Gln	D NO Gln Ile	Gly Gly Gln	Asp Asn Gln 45	Leu 30 Ser	15 Leu Gln	Gly Ala
40	Met 1 Ala	Ala Arg	(i) : (xi) Val Ala Glu 35	SEQUI () () SEQUI Val Ile 20	ENCE A) L B) T D) T UENC: Leu 5 Ala Ser	CHAI ENGT YPE: OPOL E DE Leu Val	RACT H: 2 ami OGY: SCRI Ala Gln	ERIS' 19 a no a lin PTIO Asn Lys	TICS mino cid ear N: S Leu Gly 25 Thr	aci EQ II Ala 10 Ser Gln	D NO Gln Ile	Gly Gly Gln	Asp Asn Gln 45	Leu 30 Ser	15 Leu Gln	Gly Ala
40	Met 1 Ala Phe	Ala Arg Leu Leu 50	(i): (xi) Val Ala Glu 35 Leu	SEQUION () () () () () () () () () () () () ()	ENCE A) L B) T D) T UENC: 5 Ala Ser	CHAMPERST COPOLIC CONTROL CO	RACTH: 2 ami OGY: SCRI Ala Gln Ala Asn 55	ERIS' 19 a no a lin PTIO Asn Lys Ala 40	TICS mino cid ear N: S Leu Gly 25 Thr	aci EQ I Ala 10 Ser Gln	D NO Gln Ile Phe	Gly Gln Pro 60	Asp Asn Gln 45 Xaa	Leu 30 Ser	15 Leu Gln Val	Gly Ala Asp
40	Met 1 Ala Phe Ser Met 65	Ala Arg Leu 50 Met	(i) : (xi) Val Ala Glu 35 Leu Arg	SEQUION ( ( ( ( ( ( SEQUION SE	ENCE A) L B) T D) T UENC: 5 Ala Ser Met	CHAMPERST CHAMPERST COPOLL E DE Leu Val Leu Gln Ala 70	RACTI H: 2 ami OGY: SCRI Ala Gln Ala Asn 55 Arg	ERIS' 19 a no a lin prio Asn Lys Ala 40	TICS mino cid ear N: S Leu Gly 25 Thr	aci EQ I Ala 10 Ser Gln Phe	D NO Gln Ile Phe Glu Ala 75	Gly Gln Pro 60 Leu	Asp Asn Gln 45 Xaa	Leu 30 Ser Ser	15 Leu Gln Val	Gly Ala Asp Asp 80

	Val	Leu	Phe 115	Leu	Xaa	Trp	Pro	Val 120	Met	Thr	Ala	Val	Gly 125	His	Leu	Pro
5	Pro	Pro 130	Суѕ	Val	Суѕ	Ala	Cys 135	Val	Glu	Asn	Leu	Glu 140	Thr	Asp	Cys	Cys
	Pro 145	Leu	Phe	Met	Gln	Asn 150	His	Leu	Arg	Ile	Gln 155	Phe	Thr	Leu	Cys	Cys 160
10	Pro	Ala	Ser	Pro	Leu 165	Gly	Lys	Ser	Leu	Ser 170	Cys	Phe	Ser	Leu	Leu 175	Leu
15	Pro	Pro	Pro	Leu 180	Pro	Pro	Ser	Pro	His 185	Ala	Phe	Leu	Phe	Leu 190	Val	Leu
	Thr	Leu	Leu 195	Pro	Ser	Gly	Pro	Tyr 200	Pro	Thr	Leu	Phe	Glu 205	Lys	Thr	Lys
20	Leu	Суз 210	Leu	His	Arg	Arg	Leu 215	Phe	Leu	Phe	Xaa					
	(2)	INFO	ORMAT	rion	FOR	SEO	ID N	NO: 4	115:							
25			(i) s	SEQUI	ENCE A) L	-	RACTI H: 5	ERIS 1 am	rICS ino		s				·	
30			(xi)	(	D) T	OPOL	OGY:	lin	ear	EQ II	ОИО	: 41!	5:			
	Met 1	Leu	Pro	Asp	Glu 5	Ser	Phe	Gly	Leu	Leu 10	Leu	Ser	Ile	Pro	Ser 15	Leu
35	Thr	Pro	Ser	Ala 20	Ala	Ala	Pro	Ser	Phe 25	Cys	Val	His	Leu	Met 30	Gln	Ala
40	Ser	Arg	Ser 35	Ser	Lys	Arg	Ala	Ser 40	His	Val	Pro	Val	His 45	Leu	Leu	Trp
	Gly	Asp 50	Xaa			•										
45	(2)	INFO	RMAT	CION	FOR	SEQ	ID N	io: 4	16:							
50			(i) s (xi)	() () ()	A) LI B) T O) T(	ENGTI YPE : OPOLA	H: 5 ami XGY:	0 am: no ac line	ino a cid ear	acid		: 416	<b>5</b> :			
55	Met 1	Arg	Pro	Gly	Ser 5	Phe	Ser	Phe	Ile	Ala 10	Phe	Leu	Ala	Thr	Glu 15	Val
	Ser	Ser	Cys	Phe 20	Pro	Gly	Arg	Pro	Asp 25	Суя	Xaa	Thr	Gly	Met 30	Trp	Leu
50	Leu	Gln	Leu	Gln	Lvs	Lvs	Gln	Ara	Thr	Leu	Leu	Ala	Met	Ala	Pro	Ara

	35	i	40		45
5	Arg Xaa 50				
10		TION FOR SEQ	•		
10		(A) LENGT (B) TYPE: (D) TOPOI	RACTERISTICS: TH: 70 amino a amino acid OGY: linear	acids	
15		-	SCRIPTION: SE	_	: Pro Leu Leu Leu
	1	5	Ser neu Tip	10	15
20	Leu Leu Met	Arg Leu Phe 20	Pro Leu Pro 25	Val Pro Gly	Asn Gln Arg Ala 30
	Xaa Leu Pro 35		Xaa Ala Pro 40	Arg Leu Pro	Cys Leu Leu Cys 45
25	Leu Cys Thr 50	Gln Gln Phe	Xaa Val Cys 55	Ser His Tyr 60	Leu Pro Ala Gly
30	Tyr Arg Val	Asn Ser Xaa '70			
•	(2) INFORMA	TION FOR SEQ	ID NO: 418:		
35	(i)	(A) LENGT (B) TYPE:	RACTERISTICS: H: 40 amino a amino acid OGY: linear	acids	
40			SCRIPTION: SE		
	Met His Glu 1	Lys Ala Trp 5	Asn Leu Ile	Leu Leu Trp ' 10	Trp Leu Ser Leu 15
45	Asp Leu Leu	Gly Val Ala 20	Lys Thr Ala: 25	Met Trp Ala	Gln Trp Cys Gly 30
	Leu Asn Asp	His Lys Gly	Lys Xaa 40		
50		•			,
		TION FOR SEQ			
55		(A) LENGT (B) TYPE: (D) TOPOL	RACTERISTICS: H: 22 amino a amino acid OGY: linear SCRIPTION: SE		:
50					Gln Ser Ser Xaa

```
20
                                                               15
      Gly Arg Ala Val Gln Xaa
                   20
  5
       (2) INFORMATION FOR SEQ ID NO: 420:
10
              (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 33 amins asids
                     (B) TYPE: amino aciá
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 420:
15
      Met Phe Ser Leu Leu Trp Leu Val Cys Val Pro Ser Asn Ser Ser Val
      Ala Asn Val Thr Ala Ser Arg Gly Gly Val Fhe Lys Arg Ser Leu Gly
20
                                      25
      His Glu Gly Phe Ser Xaa
               35
25
      (2) INFORMATION FOR SEQ ID NO: 421:
             (i) SEQUENCE CHARACTERISTICS:
30
                    (A) LENGTH: 35 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID MG: 421:
35
      Lys Trp Leu Leu Phe Ile Phe Leu Leu Cys Leu Gln Leu Val Asn Ala
      Leu Leu Ser Leu Phe Gln Glu Arg Phe Val His Cys Pro Ala Arg Phe
40
      Val Ser Xaa
45
      (2) INFORMATION FOR SEQ ID 10: 422:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 32 amino acids
50
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 422:
     Met Leu Leu Phe Leu Ser Ile Thr Asm Ser Leu Ser Phe Ile Ser Val
55
                5
     Asp Lys Pro Phe Gly Gln Ser Glu Asp Val Cys Pro Val Ile Ser Xaa
                  20
                          . 25
60
```

5	(2)	INFOR	ATIOI	1 FOF	SEÇ	) ID	NO:	423	:						
10		٠	) SEQI	(A) 1 (B) 1 (D) 1	LENG! IYPE IOPOI	TH: : a.m. LOGY	127 ino : li	amin acid near	o ac		D: 42	23 :			
15	Met (	Glu Ph	e Leu	Phe 5	Asn	Lys	Thr	Gly	Tr 10		Phe	≥ Ala	a Ala	Leu 15	
	Phe V	/al Le	u Ala 20	Met	Thr	Ser	Gly	Glr. 25		Trp	Asn	His	30		Gly
20	Pro I	Pro Ty 3	r Ala 5	His	Lys	Asn	Pro 40		Thr	Gly	His	Val		Туг	Ile
,	His G	Sly Se 50	r Ser	Gln	Ala	Gln 55	Phe	· Val	Ala	Glu	Thr 60		Ile	· Val	Leu
25	Leu F 65	he As	n Gly	Gly	Val 70	Thr	Leu	Gly	Met	Val 75	Leu	Leu	Cys	Glu	Ala 80
30	Ala T	hr Se	r Asp	Met 85	Asp	Ile	Gly	Lys	Arg 90	Lys	Ile	Met	Cys	Val 95	Ala
	Gly I	le Gl	y Leu 100	Val	Val	Leu	Phe	Phe 105	Ser	Trp	Met	Leu	Ser 110	Ile	Phe
35	Arg S	er Lys	s Tyr	His	Gly	Tyr	Pro 120	Tyr	Ser	Phe	Leu	Met 125	Ser	Xaa	
40	(2) I	NFORM	ATION	FOR	SEQ	ID N	10: 4	124 :							
		(i)	(	ENCE A) Li B) T	engti /PE:	H: 6: ami	9 am no a	ino . cid		s			٠	•	
45	•	(xi)	SEQ						EQ II	ONO:	: 424	1:			
	Met Th	ur Trp	His	Ser 5	Arg	Glu	Ser	Phe	Xaa 10	Leu	Leu	Arg	Val	Val 15	Ala
50	Pro Se	er Gln	Ala 20	Pro	Gly :	Met	Gln	Val 25	Ser	Pro	Ser	Gln	Arg 30	Ala	Trp
<i>55</i>	Arg Ar	g Pro	Leu	His .	Arg (	Cys	His 40	Val	Ala	Ala	Pro	Arg 45	Pro	His	His
	Phe Al	a Phe	Phe	Arg .	Asn :	Pro 55	Phe	Ser	Ттр	Ser	Phe 60	Ile	Lys	Leu	Leu
60	Tyr Ar 65	g Tyr	Leu	Xaa											

5	(2) IN	FORM?	TION	FOF	SEC	) ID	NO:	425	:						
3		(i)		JENCE (A) 1 (B) 1	LENG IYPE	TH: : am	92 a ino	mino acid	aci	ds		,			
10		(xi)	SEÇ						SEQ :	ID N	0: 4	25:	•		
	Met Gl	y Leu	Lys	Leu 5	Asn	Gly	' Arg	тут	: Ile		r Lei	ı Ile	e Lei	1 Ala	_
15	Gln Ile	≘ Ala	Tyr 20	Leu	Val	Gln	Ala	Val 25		, Ala	a Ala	a Gly	/ Lys 30		Asp
20	Ala Val	l Phe 35	Lys	Gly	Phe	Ser	Asp 40		Leu	Leu	ı Lys	Leu 45		/ Asp	Thr
	Trp Pro	j				55					60				•
25	Val His 65	: Ile	Leu	Gly	Gly 70	Phe	Pro	Gln	. Leu	His 75		His	Ser	Pro	Туr 80
	Gly Leu	Pro	Gly	Arg 85	Gly	Glu	Arg	Tyr	Val 90		Хаа				
30													•		
	(2) INF	'ORMA'	rion	FOR	SEQ	ID I	NO:	426:					•		
35		(i) :	()	A) LI B) T	ENGT YPE:	H: 3 ami	80 a no a	mino cid		.ds	•	•		-	
		(xi)		D) TY JENCI					EQ I	D NO	: 42	6:			
40	Met Ala 1	Arg	Arg	Ser 5	Ala	Phe	Pro	Ala	Ala 10	Ala	Leu	Trp	Leu	Trp 15	Ser
45	Ile Leu	Leu	Cys 20	Leu	Leu	Ala	Leu	Arg 25	Ala	Glu	Ala	Gly	Pro 30	Pro	Gln
	Glu Glu	Ser 35	Leu	Tyr	Leu	Trp	Ile 40	Asp	Ala	His	Gln	Ala 45	Arg	Val	Leu
50	Ile Gly 50	Phe	Glu	Glu .	Asp	Ile 55	Leu	Ile	Va1	Ser	Glu 60	Gly	Lys	Met	Ala
	Pro Phe 65	Thr	His .	Asp	Phe 70	Arg	Lys	Ala	Gln	Gln 75	Arg	Met	Pro	Ala	Ile 80
55	Pro Val	Asn	Ile 1	His :	Ser :	Met	Asn	Phe		Trp	Gln	Ala	Ala	Gly	Gln
				85					90					95	

•	Ile	e Met	Ala 115		Pro	Thr	Val	120		Pro	Leu	Leu	Gly 125		Val	Pro
5	His	Lys 130	Ala	Ser	Val	Val	Gln 135		Gly	Phe	Pro	Cys 140	Leu	Gly	Lys	Gln
	Asp 145	Gly	Val	Ala	Ala	Phe 150		Val	Asp	Val	Ile 155	Val	Met	Asn	Ser	Glu 160
10	Gly	Asn	Thr	Ile	Leu 165	Gln	Thr	Pro	Gln	Asn 170		Ile	Phe	Phe	Lys 175	
15	Суѕ	Gln	Gln	Ala 180	Glu	Cys	Pro	Gly	Gly 185	Cys	Arg	Asn	Gly	Gly 190	Phe	Cys
	Asn	Glu	Arg 195	Arg	Ile	Cys	Glu	Cys 200	Pro	Asp	Gly	Phe	His 205	Gly	Pro	His
20	Cys	Glu 210	Lys	Ala	Leu	Cys	Thr 215	Pro	Arg	Cys	Met	Asn 220	Gly	Gly	Leu	Cys
	Val 225	Thr	Pro	Gly	Phe	Cys 230	Ile	Cys	Pro	Pro	Gly 235	Phe	Tyr	Gly	Val	Asn 240
25	Cys	Asp	Lys	Ala	Asn 245	Суз	Ser	Thr	Thr	Суs 250	Phe	Asn	Gly	Gly	Thr 255	Суѕ
30	Phe	Tyr	Pro	Gly 260	Lys	Cys	Ile	Xaa	Pro 265	Pro	Gly	Leu	Glu	Gly 270	Glu	Gln
	Cys	Glu	Ile 275	Ser	Lys	Cys	Pro	Gln 280	Pro	Суз	Arg	Asn	Gly 285	Gly	Lys	Cys
35	Ile	Gly 290	Lys	Ser	Lys	Cys	Lys 295	Xaa	Ser	Lys	Gly	Tyr 300	Gln	Gly	Asp	Leu
	Cys 305	Ser	Lys	Pro	Val	Cys 310	Glu	Pro	Gly	Cys	Gly 315	Ala	His	Gly	Thr	Cys 320
40	His	Glu	Pro	Asn	Lys 325	Cys	Gln	Cys	Gln	Glu 330	Gly	Trp	His	Gly	Arg 335	His
45	Cys ،	Asn	Lys	Arg 340		Glu	Ala		Leu 345	Ile	His	Ala	Leu	Arg 350	Pro	Ala
	Gly	Ala	Gln 355	Leu	Arg	Gln	His	Thr 360	Pro	Ser	Leu	Lys	Lys 365	Ala	Glu	Glu
50	Arg	Arg 370	Asp	Pro	Pro	Glu	Ser 375		Tyr	Ile	Trp	Хаа 380				
55	(2)		RMAT	EQUE		CHAF	CACTE	RIST	CICS:		5					

(B) TYPE: amino acid(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 427:

```
Met Thr Ser Asn Leu Leu Leu Leu Thr Leu Leu Lys Asp Thr Leu
                                          10
     Xaa Leu Ala Lys Xaa Asn Xaa Xaa
             . 20
10
      (2) INFORMATION FOR SEQ ID NO: 428:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 47 amino acids
                    (B) TYPE: amino acid
15
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 428:
      Met Arg His His Thr Gln Leu Asn Phe Ile Phe Leu Val Glu Met Val
                                          10
20
      Phe Leu His Val Gly Gln Ala Gly Leu Lys Leu Pro Thr Ser Gly Asp
      Xaa Ala Cys Phe Gly Leu Pro Lys Val Leu Gly Leu Gln Ala Xaa
25
      (2) INFORMATION FOR SEQ ID NO: 429:
30
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 5 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
35
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 429:
      Met Cys Ser Asp Xaa
40
       (2) INFORMATION FOR SEQ ID NO: 430:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 144 amino acids
45
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 430:
       Leu Leu Ser Ile Leu Leu Cys Leu Leu Ala Ser Gly Leu Val Val Phe
50
                                          10
       Phe Leu Phe Pro His Ser Val Leu Val Asp Asp Asp Gly Ile Lys Val
                                       25
 55
       Val Lys Val Thr Phe Asn Lys Gln Asp Ser Leu Val Ile Leu Thr Ile
                                   40
       Met Ala Thr Leu Lys Ile Arg Asn Ser Asn Phe Tyr Thr Val Ala Val
 60
                              55
```

(2) INFORMATION FOR SEQ ID NO: 433:

	Thr 65	Ser	Leu	Ser	Ser	Gln 70	Ile	Gln	Tyr	Met	Asn 75	Thr	Val	Val	Asn	Phe 80
5	Thr	Gly	Lys	Ala	Glu 85	Met	Gly	Gly	Pro	Phe 90	Ser	Tyr	Val	Tyr	Phe 95	Phe
10	Cys	Thr	Val	Pro 100	Glu	Ile	Leu	Val	His 105	Asn	Ile	Val	Ile	Phe 110	Met	Arg
10	Thr	Ser	Val 115	Lys	Ile	Ser	Tyr	Ile 120	Gly	Leu	Met	Thr	Gln 125	Ser	Ser	Leu
15	Glu	Thr 130		His	Tyr	Val	Asp 135		Gly	Gly	Asn	Ser 140	Thr	Ala	Ile	Xaa
20																
	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	431:							
25					(A) 1 (B) 5 (D) 5	ENGT TYPE : TOPOI	TH: 3 : am: LOGY	TERIS	mino acid near	acio		): <b>4</b> 3	11:		•	
30	Met		Ph∈	Ph∈	Lev		· Val	. Туг	Ser	Val		Cys	Gly	Leu	Leu 15	Val
35	Тут	r Pro	Ser	: Let 20		Ser	His	s Ser	Val 25		Leu	ı Val	L Thr	: Sex		Val
	Ala	a Sei	Ala 35		ı Xaa	1										
40	(2)	) IN	FORM	AT I OI	1 FOI	R SE(	Q ID	NO:	432:	;						
45					(A) (B) (D)	LENG TYPE TOPO	TH: : am LOGY	TERI: 37 a ino : li IPTI	mino acid near	aci		0: 4	32:			
50		t Al	a Se	r Il		n Al	a Va	1 Ty:	r Ile	e Hi: 1		l Ph	e Le	u Gly	y Va: 1	l Cys 5
	Va	l Gl	n Al	a Th 2		a Al	a Cy	s Pr	o Tr		s Se	r Gl	n Cy	s Ar		a Gly
55	Se	r Va	1 Pr '3		r Xa	a										
	•															

	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 192 amino acids  (B) TYPE: amino acid															
										acı	as					
5						OPOL										
	•		(xi)	SEQU	JENC!	E DES	SCRI	PTION	N: SI	Q II	ONO:	: 433	3:			
	Met	Met	Ala	Ala	Met	Val	Leu	Thr	Ser	Leu	Ser	Cys	Ser	Pro	Val	Val
	1				5					10					15	
10	~1 <u>~</u>	Com	Dwa	Pro	C1	mh∽	C1	<b>71</b> -	n.c.	Dho	cor	בוג	Ser	λνα	λla	Δla
	GIII	261	PLO	20	GIA	1111	Giu	ALG	25	·	361	MIG	Ser	30	AIG	ALU
				•												
15	Cys	Asp	Pro 35	Trp	Lys	Glu	Ser	Gly 40	Asp	Ile	Ser	Asp	Ser 45	GIÀ	Xaa	Ser
			75													
	Thr		Ser	Gly	His	Trp		Gly	Ser	Ser	Gly		Ser	Thr	Pro	Ser
		50					55					60				
20	Pro	Pro	His	Pro	Gln	Ala	Ser	Pro	Lys	Tyr	Leu	Gly	Asp	Ala	Phe	Gly
	65					70					75					80
	Ser	Pro	Gln	Thr	Asp	His	Gly	Phe	Glu	Thr	Asp	Pro	Asp	Pro	Phe	Leu
36					85					90					95	
25	Leu	Asn	Glu	Pro	Ala	Pro	Ara	Lvs	Ara	Lvs	Asn	Ser	Val	Lvs	Val	Met
			024	100			3	-,-	105					110		
	<b>~</b>	T	<b>~</b>	Leu		Desa	<b>3</b>	~	C1	T 1 10	17-1	T OU	h-a	50×	Tle	17a l
30	Tyr	пĀS	115	Leu	πp	PIO	ASII	120	GIY	пуэ	vai	Deu	125	Jer	116	VU.
													_		•	_
	Gly	Ile 130		Arg	His	Val	Lys 135	Ala	Leu	His	Leu	G1y 140	Asp	Thr	Vai	Asp
35		Asp	Gln	Phe	Lys			Glu	Asp	Phe	Tyr 155	Tyr	Thr	Glu	Val	Gln 160
	145					150					133					100
	Leu	Lys	Glu	Glu			Ala	Ala	Ala		Ala	Ala	Ala	Ala		Pro
40					165					170					175	
	Gln	Ser	Leu	Gly	Leu	Pro	Pro	Pro	Ser	Gln	Leu	Pro	Pro	Pro	Ala	Xaa
			•	180					185					190		
45																
			•		•											
50	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	434:							
50			(i)	SEQU	ENCE	: СНА	RACT	ERIS	TICS	:						
			, -,	(	(A) I	ENG	CH: 3	31 an	nino		is					
						YPE:										
55			(xi)	SEQ		POPOI				EQ I	D NC	: 43	4:			
			•	_						_				_	_	_
	Met 1		Thr	Asn	. Туг 5		Thr	Asp	Val	Cys 10		Leu	Phe	Ser	Tyr 15	Leu
					_								·			•
60	Asn	Тут	Leu	Tyr	Phe	His	His	His	Leu	Pro	Val	Pro	Asn	Thr	Xaa	

WO 98/54963

584

	20		25	30
5	(2) INFORMATION	FOR SEO ID NO:	435:	
,		-		
10	( <i>)</i> (E (I	NCE CHARACTERI  LENGTH: 101  TYPE: amino  TOPOLOGY: 1: ENCE DESCRIPTI	amino acids acid	: 435:
15	Met Gly Phe Phe 1	Phe Val Leu Ph 5	ne Phe Leu Tyr 10	Leu Ala Leu Ser Arg 15
13	Asp Trp Ser Ile 20	Asn Phe Leu Ly	ys Asp His Arg 25	Ile Asn Phe Phe Val
20	Ala Thr Ser Tyr 35		r Val Arg Gly 10	Xaa Pro Xaa Val Pro 45
	Ala Asp Thr Pro	Leu Gly Pro Le 55	eu Leu Ser Leu	Trp Leu His His Asn 60
25	Ala Phe Phe Ser 65	Ile Leu Pro Ly 70	ys Phe Pro Glu 75	Asn Xaa Xaa Phe Leu 80
20	Ile Leu Lys Lys	Leu Val Val G	lu Met Gly Trp 90	Asp Leu Phe Ile Ser 95
30	Pro Glu Asn Lys 100	Xaa	:	
35	(2) INFORMATION	FOR SEQ ID NO	: 436:	
40	(	ENCE CHARACTER A) LENGTH: 37 B) TYPE: amino D) TOPOLOGY: 1 UENCE DESCRIPT	amino acids acid	o: <b>4</b> 36:
45	Met Ala Arg Tyr 1	Phe Ile Phe P	he Ile Leu Val 10	Phe Met Lys Val Ser 15
	Leu Asn Thr Thr	Trp Pro Ala P	ro Arg Pro Ala 25	Thr Leu Arg Thr Ala 30
50	Asn Lys Ser Lys	Xaa		
55	(2) INFORMATION	FOR SEQ ID NO	): <b>43</b> 7:	

(i) SEQUENCE CHARACTERISTICS:

60

(A) LENGTH: 42 amino acids
(B) TYPE: amino acid

(D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 437:													
5	Phe Ser Thr Ile Arg Ser Gly Leu Thr Asp Arg Ser Val Asn Phe Leu  1 5 10 15													
J	Phe Leu Phe Leu Asp Val Pro Asp Cys Arg Leu Val Asn Ile Glu Leu 20 25 30													
10	Met Ala Asn Ser Thr Val Thr His Ala Xaa 35 40													
15	(2) INFORMATION FOR SEQ ID NO: 438:  (i) SEQUENCE CHARACTERISTICS:													
	<ul><li>(A) LENGTH: 1 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>													
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 438:													
	Leu 1													
25														
	(2) INFORMATION FOR SEQ ID NO: 439:													
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 25 amino acids													
	(B) TYPE: amino acid (D) TOPOLOGY: linear													
0.5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 439:													
35	Met Pro Trp Arg Arg Ala Gly Leu Met Met Leu Pro Ile Ile Thr Gly 1 5 10 15													
	Cys Cys Pro Cys Ser Ala Ser Ile Xaa 20 25													
40														
	(2) INFORMATION FOR SEQ ID NO: 440:													
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 54 amino acids  (B) TYPE: amino acid													
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 440:													
50	Met Tyr Leu Cys Lys Thr Val Lys Val Leu Ile Cys Tyr Asp Trp Ile 1 5 10 15													
<b>5</b> 5	Leu Gly Leu Val Ser Ser Gly Gln His Trp Val Val Ser Leu Ser Tyr 20 25 30													
	Ser Ile Arg Val Tyr Pro Ala Met His Phe Thr Leu Cys Val His Ile 35 40 45													
60	Tyr Ser Lys Glu Pro Cys													

PCT/US98/11422

5	(2) INFORMATION FOR SEQ ID NO: 441:
10	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 42 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 441:</li> </ul>
15	Met Thr Ala Leu Val Trp Arg Lys Gly Pro Asp Gly Gly Ser Arg Lys  1 5 10 15
13	Pro Ile Leu Leu Phe Phe Phe Leu Pro Leu Ile Leu Cys Phe His 20 25 30
20	Ser Phe Ile His Ser Ser Asn Ile Cys Xaa 35 40
25	(2) INFORMATION FOR SEQ ID NO: 442:
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 66 amino acids  (B) TYPE: amino acid
30	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 442:
	Met Phe Leu Thr Trp Phe Leu Leu Leu Ser Val Ala Trp Xaa Ala 1 5 10 15
35	Leu Thr Arg Ser Gly Arg Ser Cys Leu Pro Leu Val Gly Arg Pro Arg 20 25 30
40	Glu Gln Ser Pro Arg Thr His Cys Ala Ala Ser Ser Thr Lys Glu Arg 35 40 45
	Asn Ser Asp Pro Gln Pro Ser Pro Pro Glu Val Val Gly Pro Leu Trp 50 .55 60
45	Ser Xaa 65
50	(2) INFORMATION FOR SEQ ID NO: 443:
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 156 amino acids  (B) TYPE: amino acid
55	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 443:
	Met Lys Ala Ile Gly Ile Glu Pro Ser Leu Ala Thr Tyr His His Ile 1 5 10 15
60	Ile Arg Leu Phe Asp Gln Pro Gly Asp Pro Leu Lys Arg Ser Ser Phe

				20					25					30		
	Ile	Ile	Туг 35	Asp	Ile	Met	Asn	Glu 40	Leu	Met	Gly	Lys	Arg 45	Phe	Ser	Pro
5	Lys	Asp 50	Pro	Asp	Asp	Asp	Lys 55	Phe	Phe	Gln	Ser	Ala 60	Met	Ser	Ile	Cys
10	Ser 65	Ser	Leu	Arg	Asp	Leu 70	Glu	Leu	Ala	Tyr	Gln 75	Val	His	Gly	Leu	Leu 80
	Lys	Thr	Gly	Asp	Asn 85	Trp	Lys	Phe	Ile	Gly 90	Pro	Asp	Gln	His	Arg 95	Asn
15	Phe	Tyr	Tyr	Ser 100	Lys	Phe	Phe	Asp	Leu 105	Ile	Cys	Leu	Met	Glu 110	Gln	Ile
20	Asp	Val	Thr 115	Leu	Lys	Trp	Tyr	Glu 120	Asp	Leu	Ile	Pro	Ser 125	Ala	Tyr	Phe
20	Pro	His 130		Gln	Thr	Met	Ile 135		Leu	Leu	Gln	Ala 140	Leu	Asp	Val	Ala
25	Asn 145	Arg	Leu	Glu	Val	Ile 150	Pro	Lys	Ile	Trp	Glu 155					
30	(2)	INF			ENCE		RACT	TERIS	TICS		ls					
35			(xi)		(D) :	ropoi	LOGY	: li	near	SEQ I	D NO	o: 44	14:			
	Met 1		Ph∈	e Leu	Phe 5		Phe	e Ile	e Val	Phe 10		з Тут	Leu	Tr	Gly 15	Leu
40	Phe	Thi	Ala	20 20		g Glr	Lys	s Lys	Glu 25		sei	Thr	: Glu	Glu 30		. Lys
45			39 Let					4( 5 Let	)		s Sei	c Lys	Thi 45		. Lys	: Lys
50	(2)			ATIO	1 FO	R SE(			445	:						
55		٠		SEQ	(A) (B) (D)	LENG TYPE TOPO	TH: : an LOGY	416 ino : li	amin acid near	o ac l		io: 4	45:			
60		t Ar 1	g Th	r Le		e As: 5	n Le	u Le	u Tr	p Le		a Le	u Al	a Cy	s Se	r Pro 5

	Val	His	Thr	Thr 20	Leu	Ser	Lys	Ser	Asp 25	Ala	Lys	Lys	Ala	Ala 30	Ser	Lys
5	Thr	Leu	Leu 35	Glu	Lys	Ser	Gln	Phe 40	Ser	Asp	Lys	Pro	Val 45	Gln	Asp	Arg
10	Gly	Leu 50	Val	Val	Thr	Asp	Leu 55	Lys	Ala	Glu	Ser	Val 60	Val	Leu	Glu	His
10	Arg 65	Ser	Tyr	Cys	Ser	Ala 70	Lys	Ala	Arg	Asp	Arg 75	His	Phe	Ala	Gly	Asp 80
15	Val	Leu	Gly	Tyr	Val 85	Thr	Pro	Trp	Asn	Ser 90	His	Gly	Tyr	Asp	Val 95	Thr
	Lys	Val	Phe	Gly 100	Ser	Lys	Phe	Thr	Gln 105	Ile	Ser	Pro	Val	Trp 110	Leu	Gln
20	Leu	Lys	Arg 115	Arg	Gly	Arg	Glu	Met 120	Phe	Glu	Val	Thr	Gly 125	Leu	His	Asp
25	Val	Asp 130		Gly	Trp	Met	Arg 135	Ala	Val	Arg	Lys	His 140	Ala	Lys	Gly	Leu
23,	His 145		Val	Pro	Arg	Leu 150	Leu	Phe	Glu	Asp	Trp 155	Thr	Tyr	Asp	Asp	Phe 160
30	Arg	Asn	Val	Leu	Asp 165	Ser	Glu	Asp	Glu	Ile 170		Glu	Leu	Ser	Lys 175	
	Val	Val	Gln	Val 180		Lys	Asn	Gln	His 185		Asp	Gly	Phe	Val 190		Glu
35	Val	. Trp	Asn 195		Leu	Leu	Ser	Gln 200		Arg	Val	Gly	Leu 205		His	Met
40	Leu	210		Leu	ı Ala	Glu	Ala 215		His	Gln	Ala	Arg 220		. Leu	Ala	Leu
	Let 225		. Ile	Pro	Pro	Ala 230		• Thr	Pro	Gly	7 Thr 235		Glr	. Leu	Gly	Met 240
45	Phe	e Thi	: His	Lys	Glu 245		: Glu	ı Glm	Leu	Ala 250		Val	Lev	ı Asp	Gly 255	Phe
	Sei	c Let	ı Met	Thu 260		Ası	тул	Ser	Thr 265		a His	Glr	ı Pro	Gly 27(		Asn
50	Ala	a Pro	275		Tr	Va:	LAr	y Ala 280		val	l Glr	ı Val	L Let 289		Pro	Lys
55	Se	r Ly: 29		) Ar	g Sei	. Ly:	29!		ı Lev	ı Gly	y Let	1 Asi 300		е Тут	c Gly	/ Met
<i>JJ</i>	As <sub>3</sub>		r Ala	a Thi	r Sei	31		p Ala	a Arq	g Gl	u Pro 319		l Va	l Gly	/ Ala	a Arg 320
60	ТУ	ż Il	e Gli	n Th	r Lei		s As	p His	s Ar	g Pro		g Me	t Va	l Tr	As <sub>1</sub>	o Ser

	Gln X	aa	Ser	Glu 340	His	Phe	Phe	Glu	Tyr 345	Lys	Lys	Ser	Arg	Ser 350	Gly	Arg
5	His V		Val 355	Phe	Tyr	Pro	Thr	Leu 360	Lys	Ser	Leu	Gln	Val 365	Arg	Leu	Glu
10	Leu A	Ala 370	Arg	Glu	Leu	Gly	Val 375	Gly	Val	Ser	Ile	Trp 380	Glu	Leu	Ala	Arg
10	Ala 1 385	(rp	Thr	Thr	Ser	Thr 390	Thr	Суѕ	Ser	Arg	Trp 395	Ala	Leu	Arg	Pro	Pro 400
15	Arg 1	Crp	Thr	Cys	Ser 405	Phe	Leu	Ser	His	Gly 410	Val	Ser	Glu	Gln	Val 415	Xaa
20																
	(2)					SEQ CHA				:						
25	•			(	(A) I (B) I (D) I	ENGI YPE: YPOI E DE	H: 6 ims :YDO.	54 an ino a : lir	mino acid near	ació		): <b>4</b> 4	6:			
30	Met .	Ala	Pro	Gly	Pro		Ser	Ala	Thr	Gln 10		Val	Val	Ile	His 15	Thr
35	Thr	His	Cys	Leu 20		Leu	Pro	Val	. Trp 25		. Leu	Ser	Leu	Val 30		Glu
55	Leu	Leu	Gly 35		, Ala	Pro	Pro	His 40		Lys	: Asp	Ala	. Leu 49		pro	Ser
40	Lys	Lys 50		. Lys	: Lys	. Lys	Let 55		a Gly	, Gl	Pro	Va]		) Ile	Pro	Pro
45																
	(2)	INF	ORM	OITA	1 FOI	R SE(	O ID	NO:	447	:						
50					(A) (B) (D)	E CHI LENG TYPE TOPO CE D	TH: : an LOGY	206 ino ': li	amin acid near	o ac		O: 4	47:			
55	Met 1		ı Gl	y Ala		s Pro	o Hi	s Tr	p Le	u Pr		y Pr	o Le	u Hi	s Se 1	r Pro 5
	Gly	Le	ı Pr	o Le		l Le	u Va	l Le	u Le 2		a Le	u Gl	y Al	a Gl		p Ala
60																

	Gln	Glu	Gly 35	Ser	Glu	Pro	Val	Leu 40	Leu	Glu	Gly	Glu	Cys 45	Leu	Val	Val
5	Cys	Glu 50	Pro	Gly	Arg	Ala	Ala 55	Ala	Gly	Gly	Pro	Gly 60	Gly	Ala	Ala	Leu
	Gly 65	Glu	Ala	Pro	Pro	Gly 70	Arg	Val	Ala	Phe	Ala 75	Ala	Val	Arg	Ser	Xaa 80
10	His	His	Glu	Pro	Ala 85	Gly	Glu	Thr	Gly	Asn 90	Gly	Thr	Xaa	Gly	Ala 95	Ile
15	Tyr	Phe	Asp	Gln 100	Val	Leu	Val	Asn	Glu 105	Gly	Gly	Gly	Phe	Asp 110	Arg	Ala
13	Ser	Gly	Ser 115	Phe	Val	Ala	Pro	Val 120	Arg	Gly	Val	Tyr	Ser 125	Phe	Arg	Phe
20	His	Val 130		Lys	Val	Tyr	Asn 135	Arg	Gln	Thr	Val	Gln 140		Ser	Leu	Met
	Leu 145	Asn	Thr	Trp	Pro	Val 150		Ser	Ala	Phe	Ala 155	Asn	Asp	Pro	Asp	Val 160
25	Thr	Arg	Glu	Ala	Ala 165		Ser	Ser	Val	Leu 170		Pro	Leu	Asp	Pro 175	Gly
30	Asp	Arg	Val	Ser 180		Arg	Leu	Arg	Arg 185		Asn	Leu	Leu	Gly 190		Trp
	Lys	Тут	Ser 195		Phe	Ser	Gly	Phe 200		Ile	Phe	Pro	205			
35	(2)	INE	ORMA	<b>MOITA</b>	1 FOF	SEÇ	ΩÍD	NO:	448:							
40				SEQI	(A) : (B) : (D) :	LENG TYPE TOPO	TH: : am LOGY	62 a ino a : li:	mino acid near	aci		o: 4	48:			
45	Met 1		r Sei	c Le		ı Sei	c Ala	a Gly	/ Let	1 Gl: 10		a Se	r Lei	ı Cys	Gl <sub>3</sub>	/ Lys
	Xaa	a Le	ı Tr	p Ala 2		r Th	r Trį	у Туг	Let 2!		l Cy:	s Cy:	s Le	Leu 30		) Phe
50	Phe	e Hi	s Gl		у Су:	s Cy	s As	p Hi:		s Se	r Ly:	s Gl	n Gli 4		r Il	e Pro
55	Ası		u Ly 0	s Se	r Ty	r Cy	s Gl		u Se:	r Th	r Il	e Gl 6	u Il O	e Xa	a	
	(2	) IN	FORM	ATIO	n fo	R SE	Q ID	NO:	449	:						

(i) SEQUENCE CHARACTERISTICS:

				(E	3) T'S	NGTH PE:	amir	no ac	id	acic	11.5					
_			(xi)	SEQU		POLC DES				Q II	NO:	449	:			
5	Met 1	Ser	Thr	Lys	Lys 5	Leu	Cys	Ile '	Val (	Gly 10	Gly	Ile	Leu	Leu	Val 15	Phe
10	Gln	Ile	Ile	Ala 20	Phe	Leu	Val	Gly	Gly 25	Leu	Ile	Ala	Pro	Gly 30	Pro	Thr
	Thr	Ala	Val 35	Ser	Tyr	Met	Ser	Val 40	Lys	Cys	Val	Asp	Ala 45	Arg	Lys	Asn
15	His	His 50	Lys	Thr	Lys	Trp	Phe 55	Val	Pro	Trp	Gly	Pro 60	Asn	His	Cys	Asp
20	Lys 65	Ile	Arg	Asp	Ile	Glu 70	Glu	Ala	Ile	Pro	Arg 75	Glu	Ile	Glu	Ala	Asn 80
				Phe	85					90					95	
25				Gln 100					105					110		•
			115					120					125			
30		130		Tyr			135					140				
35	145			Val		150			•		155					160
				His	165					170					175	
40				180					185					190		
			195	5				200					205	i		Glu 
45		210	)				215					220				Thr
50	225	5				230					235			•		240
					245	5				250	)				255	
55				260	)				265	5				270		Phe
			27	5			•	280	)				289	5		Trp
60	Me	t Le	u Le	u Phe	e Gly	y Ast	Ile	e Arg	, Glr	n Ala	a Ser	: Sei	Me	t Xaa	. Cys	s Phe

```
300
                              295
         290
     Xaa Pro Ser Gly Ser Ser Ser Val Ala Ser Thr Xaa
                         310
5
      (2) INFORMATION FOR SEQ ID NO: 450:
10
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 24 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 450:
15
      Met Leu Ala Leu Leu Gly Leu Leu Ala Gly Thr Glu His Pro Pro Gly
                                        10
      Pro Gln Gly Pro Gly Pro Ser Xaa
20
                   20
      (2) INFORMATION FOR SEQ ID NO: 451:
25
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 10 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
30
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 451:
      Met Pro Ser Gly Ala Cys Cys Ser Pro Xaa
                        5
35
       (2) INFORMATION FOR SEQ IQ NO: 452:
              (i) SEQUENCE CHARACTERISTICS:
40
                     (A) LENGTH: 26 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 452:
       Met Leu Pro Ala Leu Ser Thr Val Leu Leu Pro Thr Pro Ser Leu Cys
45
                        5
       Ser Gly Asn Pro Arg Glu Gly Trp Ala Xaa
                    20 .
 50
       (2) INFORMATION FOR SEQ ID NO: 453:
 55
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 172 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
               (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 453:
 60
```

	Met 1	Tyr	Ser	Leu	His 5	Ser	Trp	Val	Gly	Leu 10	Ile	Ala	Val	Ile	Cys 15	Tyr
5	Leu	Leu	Gln	Leu 20	Leu	Ser	Gly	Phe	Ser 25	Val	Phe	Leu	Leu	Pro 30	Trp	Ala
	Pro	Leu	Ser 35	Leu	Arg	Ala	Phe	Leu 40	Met	Pro	Ile	His	Val 45	Tyr	Ser	Gly
10	Ile	Val 50	Ile	Phe	Gly	Thr	Val 55	İle	Ala	Thr	Ala	Leu 60	Met	Gly	Leu	Thr
15	Glu 65	Lys <sub>.</sub>	Leu	Ile	Phe	Ser 70	Leu	Arg	Asp	Pro	<b>Ala</b> 75	Tyr	Ser	Thr	Phe	Pro 80
	Pro	Glu	Gly	Val	Phe 85	Val	Asn	Thr	Leu	Glý 90	Leu	Leu	Ile	Leu	Val 95	Phe
20	Gly	Ala	Leu	Ile 100	Phe	Trp	Ile	Val	Thr 105	Arg	Pro	Gln	Trp	Lys 110	Arg	Pro
	Lys	Glu	Pro 115	Asn	Ser	Thr	Ile	Leu 120	His	Pro	Asn	Gly	Gly 125	Thr	Glu	Gln
25	Gly	Ala 130	Arg	Gly	Ser	Met	Pro 135	Ala	Tyr	Ser	Gly	Asn 140	Asn	Met	Asp	Lys
30	Ser 145	Asp	Ser	Glu	Leu	<b>A</b> sn 150	Xaa	Glu	Val	Ala	Ala 155	Arg	Lys	Arg	Asn	Leu 160
	Ala	Leu	Asp	Glu	Ala 165		Gln	Arg	Ser	Thr 170	Met	Xaa				
35	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	454:							
40			•	1	(A) I (B) T (D) T	ENGT TYPE : TOPOI	TH: 9 ami	6 an ino a : lir	nino acid near	: acid		): 45	4:			
45	Met 1		His	Val	Leu 5		Ala	Gln	Val	Thr 10		Val	Ile	Ile	Thr 15	Thr
	Val	. Ser	Val	. Leu 20		Phe	asp	Phe	Arg 25		Ser	Leu	Glu	Phe 30		Leu
50	Glu	. Ala	Хаа 35		· Val	. Xaa	. Leu	Ser 40		Phe	·Ile	туг	Asn 45		Ser	Lys
55	Pro	Glr 50		. Pro	Glu	Тут	: Ala		Arg	Gln	Glu	Arg 60		Arg	Asp	Leu
<i></i>	Ser 65		/ Asr	ı Lev	Tr	70		, Sex	: Ser	Gly	Asr 75		Glu	Glu	. Leu	Glu 80
60 .	Arg	J Leu	ı Thi	: Lys	Pro 85		s Ser	: Asr	Glu	Ser 90		Glu	a Asp	Thr	Phe 95	Xaa

5	
	(2) INFORMATION FOR SEQ ID NO: 455:
10	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 171 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 455:</li> </ul>
15	Met Arg Gly Pro Ala Gln Ala Lys Leu Leu Pro Gly Ser Ala Ile Gln 1 5 10 15
20	Ala Leu Val Gly Leu Ala Arg Pro Leu Val Leu Ala Leu Leu Leu Val 20 25 30
20	Ser Ala Ala Leu Ser Ser Val Val Ser Arg Thr Asp Ser Pro Ser Pro 35 40 45
25	Thr Val Leu Asn Ser His Ile Ser Thr Pro Asn Val Asn Ala Leu Thr 50 55 60
	His Glu Asn Gln Thr Lys Pro Ser Ile Ser Gln Ile Ser Thr Thr Leu 65 70 75 80
30	Pro Pro Thr Thr Ser Thr Lys Lys Ser Gly Gly Ala Ser Val Val Pro 85 90 95
35	His Pro Ser Pro Thr Pro Leu Ser Gln Glu Glu Ala Asp Asn Asn Glu 100 105 110
55	Asp Pro Ser Ile Glu Glu Glu Asp Leu Leu Met Leu Asn Ser Ser Pro 115 120 125
40	Ser Thr Ala Lys Asp Thr Leu Asp Asn Gly Asp Tyr Gly Glu Pro Asp 130 135 140
	Tyr Asp Trp Thr Thr Gly Pro Arg Asp Asp Glu Ser Asp Xaa His 145 150 155 160
45	Leu Gly Arg Lys Gln Gly Leu His Gly Asn Xaa 165 170
50	(2) INFORMATION FOR SEQ ID NO: 456:
55	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 92 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 456:
	Met Lys Ala Ser Gln Cys Cys Cys Cys Leu Ser His Leu Leu Ala Ser

	Val	Leu	Leu	Leu 20	Leu	Leu	Leu	Pro	Glu 25	Leu	Ser	Gly	Xaa	Leu 30	Xaa	Val
5	Leu	Leu	Gln 35	Ala	Ala	Glu	Ala	Ala 40	Pro	Gly	Xaa	Gly	Pro 45	Pro	Asp	Pro
	Arg	Pro 50	Gly	His	Tyr	Arg	Arg 55	Cys	His	Arg	Ala	Leu 60	Thr	Pro	Ala	Gln
10	Gln 65	Pro	Gly	Arg	Gly	Leu 70	Ala	Glu	Ala	Ala	Gly 75	Ala	Ala	Gly	Leu	Arg 80
15	Gly	Arg	Gln	Trp	Gln 85	Gln	Pro	Cys	Gly	Arg 90	Ala	Xaa				
20	(2)	INF		SEQU )	FOR ENCE A) L B) T D) T	CHA ENGT YPE:	RACT H: 2 ami	ERIS 06 a no a	TICS mino cid		ds ·					
25	Ile 1	Ser	•	SEQ	UENC	E DE	SCRI	PTIO	N: S					Leu	Pro 15	Glu
30		Thr	Ala	Glu 20	Ser	Leu	Glu	Ala	Gly 25	Asp	Ser	Asn	Gln	Phe 30	Cys	Trp
	Arg	Asn	Leu 35		Ser	Cys	Ile	Asn 40	Leu	Leu	Arg	Ile	Leu 45		Lys	Leu
35	Thr	Lys 50		Lys	His	Ser	Arg 55		Met	Met	Leu	Val 60	Val	Phe	Lys	Ser
40	Ala 65		Ile	e Leu	Lys	Arg 70		Leu	Lys	Val	Lys 75		Ala	Met	Met	Gln 80
	Leu	Тут	· Val	. Leu	Lys 85		Leu	Lys	Val	Gl::		Lys	Tyr	Leu	Gly 95	Arg
45	Gln	Trp	Arg	100		Asn	Met	. Lys	105		: Ser	Ala	Ile	110		Lys
	Val	. Arg	His 115		, Leu	Asn	Asp	120		Ala	тут	Gly	125		Leu	Asp
50	Ala	130		Tr	Asp	Phe	Glr 139		Glu	ı Glu	ı Cys	140		ı Arg	, Ala	Asn
55	11e		ı Arç	g Phe	e Asr	150		y Arg	у Туг	: Asg	155		His	s Ser	Asr	160
	Asg	Phe	e Le	ı Pro	Val 165		) Ası	ı Cys	: Lev	170		. Val	. Le	ı Gly	/ Glr 175	Arg
60	Va]	L Asy	o Le	u Pro		ı Ası	Phe	e Glr	n Met 185		а Туг	: Asp	Le	1 Trg		ı Glu

	Arg Glu Va 19		Lys Pro	Ile Ser ' 200	Trp Glu Glu	Leu Leu 205
5		•				
	(2) INFORM	MATION FOR	SEQ ID	NO: 458:		
10		(A) 1 (B) ( (D) (	LENGTH: 3 TYPE: ami	: linear		B:
15	Met Ala Pr	o Pro Ala		Pro Ala	Ser Gly Gly 10	Ser Gly Glu Val 15
20	Asp Glu Le	eu Phe Asp 20	Val Lys	Asn Ala 25	Phe Tyr Ile	Gly Ser Tyr Gln 30
20		le Asn Glu 35	ı Ala Xaa	a Xaa Val 40	Lys Leu Ser	Ser Pro Glu Arg 45
25	Asp Val G	lu Arg Ası	Val Phe		Arg Ala Tyr 60	Leu Ala Gln Arg
	Lys Phe G	ly Val Va	Leu Asr 70	o Glu Ile	Lys Pro Ser 75	Ser Ala Pro Glu 80
30	Leu Gln A	la Val Arg		e Ala Asp	Tyr Leu Ala 90	His Glu Ser Arg 95
35	Arg Asp S	er Ile Va 100	l Ala Gl	u Leu Asp 105	Arg Glu Met	Ser Arg Ser Xaa 110
55		hr Asn Th 15	r Thr Ph	e Leu Leu 120	Met Ala Ala	Ser Ile Tyr Leu 125
40	His Asp G 130	ln Asn Pr	o Asp Al		Arg Ala Leu 140	His Gln Gly Asp
	Ser Leu G 145	lu Cys Th	r Ala Me 150	t Thr Val	Gln Ile Leu 155	Leu Lys Leu Asp 160
45	Arg Leu A	sp Leu Al 16		s Glu Leu	Lys Arg Met 170	: Gln Asp Leu Asp 175
50	Glu Asp A	la Thr Le 180	u Thr Gl	n Leu Ala 185		Val Ser Leu Ala 190
30		Gly Glu Ly 195	s Leu Gl	n Asp Ala 200	Tyr Tyr Ile	e Phe Gln Glu Met 205
55	Ala Asp I 210	Jys Cys S€	er Pro Th		Leu Leu Ass 220	n Gly Gln Ala Ala )
	Cys His N 225	Met Ala Gl	n Gly Ar 230	g Trp Glu	Ala Ala Glu 235	u Gly Leu Leu Gln 240
60	Glu Ala I	Leu Asp Ly	s Asp Se	er Gly Tyn	Pro Glu Th	r Leu Val Asn Leu

			:	245					250				:	255	
5	Ile Val		Ser ( 260	Gln I	His	Leu		Lys 265	Pro	Pro	Glu '		Thr 2 270	Asn i	Arg
-5	Tyr Leu	Ser 275	Gln :	Leu	Lys	Asp	Ala 280	His	Arg	Ser		Pro 285	Phe	Ile	Lys
10	Glu Tyr 290		Ala	Lys	Glu	Asn 295	Asp	Phe	Asp	Arg	Leu 300	Val	Leu	Gln	Tyr
	Ala Pro 305	Ser	Ala	Glu	Ala 310	Gly	Pro	Glu	Leu	Ser 315	Gly	Pro			
15															
	(2) INF														
20			()	A) L B) T D) T	ENGT YPE: OPOL	H: 2 ami OGY:	61 a no a lin	mino cid ear	aci		45	•			
			SEQ												
25	Arg Ası	Val	Glu	Arg 5	Asp	Val	Phe	Leu	Tyr 10	Arg	Ala	Tyr	Leu	Ala 15	Gln
30	Arg Ly	s Phe	Gly 20		Val	Leu	Asp	Glu 25	Ile	Lys	Pro	Ser	Ser 30	Ala	Pro
30	Glu Le	u Gln 35		Val	Arg	Met	Phe 40		Asp	Tyr	Leu	Ala 45	His	Glu	Ser
35	Arg Ar		Ser	Ile	Val	Ala 55		Leu	Asp	Aṛg	Glu 60		Ser	Arg	Ser
	Xaa As 65	p Val	. Thr	Asn	Thr 70		Phe	. Leu	. Leu	Met 75		Ala	Ser	Ile	Tyr 80
40	Leu Hi	s Asp	Gln	Asn 85		Asp	Ala	Ala	Leu 90		Ala	Leu	His	Gln 95	Gly
45	Asp Se	r Lev	100		Thi	c Ala	Met	105		Glr	ıle	Leu	110	Lys	Leu
43	Asp Ar	g Let 119		Leu	ı <sub>.</sub> Ala	a Arg	120		ı Lev	ı Lys	Arg	125	Gln	. Asp	Leu
50	Asp Gl		Ala ç	a Thr	. Le	u Thi 139		ı Lev	ı Ala	a Thu	Ala 140		Val	. Ser	Leu
	Ala Tì 145	ır Gl	y Gly	y Glu	15:		u <sub>.</sub> Glı	n Ası	o Ala	a Tyr 15		: Ile	e Phe	Glr	160
55	Met A	la As	p Ly:	s Cy:	_	r Pr	o Th	r Lei	ı Le		ı Le	ı Ası	ı Gly	7 Glr 175	n Ala S
60	Ala C	ys Hi	s Me		a Gl	n Gl	y Ar	g Trj 18		u Al	a Ala	a Glı	190	/ Let	ı Leu

•	Gln	Glu	Ala 195	Leu	Asp	Lys	Asp	Ser 200	Gly	Tyr	Pro	Glu	Thr 205		Val	Asn
5		Ile 210	Val	Leu	Ser	Gln	His 215	Leu	Gly	Lys	Pro	Pro 220	Glu	Val	Thr	Asn
	Arg 225	Tyr	Leu	Ser	Gln	Leu 230	Lys	Asp	Ala	His	Arg 235	Ser	His	Pro	Phe	Ile 240
10	Lys	Glu	Tyr	Gln	Ala 245	Lys	Glu	Asn	Asp	Phe 250	Asp	Arg	Leu	Val	Leu 255	Gln
15	Tyr	Ala	Pro	Ser 260	Ala										ı	
								٠								
	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	. : ON	460:							
20			(i)			CHA ENGT					ds		•			
				(	D) T	YPE: OPOL	OGY:	lin	ear			4.0	•		•	
25						E DE								•••	**! =	71.
	Met 1		Ala	Ile	Gly 5	Ile	Glu	Pro	Ser	Leu 10	Ala	Thr	ıyr	Hls	н1S	11e
30	Ile	Arg	Leu	Phe 20	Asp	Gln	Pro	Gly	Asp 25		Leu	Lys	Arg	Ser 30	Ser	Phe
	Ile	Ile	Tyr 35		Ile	Met	Asn	Glu 40		Met	Gly	Lys	Arg 45		Ser	Pro
35	Lys	Asp 50		Asp	Asp	Asp	Lys 55		Phe	Gln	Ser	Ala 60		Ser	Ile	Суз
40	Ser 65		Leu	Arg	Asp	Leu 70		Leu	Ala	Tyr	Gln 75		. His	Gly	Leu	. Leu 80
	Lys	Thr	Gly	Asp	Asn 85		Lys	Phe	: Ile	Gly 90		Asp	Gln	His	Arg 95	Asn
45	Phe	тут	Туг	Ser 100		Phe	Phe	Asp	105		Cys	Leu	Met	: Glu 110		lle
	Asp	Val	115		Lys	Trp	Туг	120		Leu	Ile	Pro	125		тут	Phe
50	Pro	130		Glr	Thr	Met	135		: Lev	. Leu	Glr	140		ı Asp	Va]	. Ala
55	Asr 145		g Leu	ı Glu	Va]	l Il∈ 150		Lys	Ile	Trţ	Glu 159					
	(2)	IN	FORM	TOITE	I FOI	R SEC	) ID	NO:	461:	:						

(i) SEQUENCE CHARACTERISTICS:

PCT/US98/11422

				(I (I	3) TY	PE:	amir XGY:	76 ar no ac line	cid ear							
•		(	(xi)	SEQU	ENCE	DES	CRIE	OITS	1: SE	Q II	ONO:	461	.:			
5	Lys i	Asp	Ser	Lys	Glu 5	Tyr	Gly	His	Thr	Phe 10	Arg	Ser	Asp	Leu	Arg 15	Glu
10	Glu :	Ile		Met 20	Leu	Met	Ala	Arg	Asp 25	Lys	His	Pro	Pro	Glu 30	Leu	Gln
	Val .	Ala	Phe 35	Ala	Asp	Cys	Ala	Ala 40	Asp	Ile	Lys	Ser	Ala 45	Tyr	Glu	Ser
15	Gln	Pro 50	Ile	Arg	Gln	Thr	Ala 55	Gln	Asp	Trp	Pro	Ala 60	Thr	Ser	Leu	Asn
20	Cys 65	Ile	Ala	Ile	Leu	Phe 70	Leu	Arg	Ala	Gly	Arg 75	Thr	Gln	Glu	Ala	Trp 80
20	Lys	Met	Leu	Gly	Leu 85	Phe	Arg	Lys	His	Asn 90	Lys	Ile	Pro	Arg	Ser 95	Glu
25	Leu	Leu	Asn	Glu 100	Leu	Met	Asp	Ser	Ala 105	Lys	Val	Ser	Asn	Ser 110	Pro	Ser
	Gln	Ala	Ile 115	Glu	Val	Val	Glu	Leu 120	Ala	Ser	Ala	Phe	Ser 125	Leu	Pro	Ile
30	Cys	Glu 130		Leu	Thr	Gln	Arg 135	Val	Met	Ser	Asp	Phe 140	Ala	Ile	Asn	Gln
35	Glu 145	Gln	Lys	Glu	Ala	Leu 150		Asn	Leu	Thr	Ala 155	Leu	Thr	Ser	Asp	Ser 160
	Asp	Thr	Asp	Ser	Ser 165		Asp	Ser	Asp	Ser 170	Asp	Thr	Ser	Glu	Gly 175	
40										-						
45	(2)	INF						NO:								
			(1)	_	(A) 1 (B) 1	LENG TYPE	rh: :	TERIS 324 a ino a	amino acid		ids		-			
50			(xi)					: li		SEQ :	ID NO	): 46	52:			
	Met 1		Ser	Asr	Asr S		ser	: Asp	Il∈	Glu 10		Glu	a Asg	Let	1 Lys	Leu
55	Glu	Let	ı Arg	arg 20		ı Arg	J Ası	Lys	His 25		ı Lys	Glu	ı Ile	e Glr 30		Leu
60	Gln	Sei	c Arg		ı Lys	s His	s Glı	1 Ile 40		ı Sei	c Lev	туз	Th:		s Ée	ı. Gly.

	Lys	Val 50	Pro	Pro	Ala	Val	Ile 55	Ile	Pro	Pro	Ala	Ala 60	Pro	Leu	Ser	Gly
5	Arg 65	Arg	Arg	Arg	Pro	Thr 70	Lys	Ser	Lys	Gly	Ser 75	Lys	Ser	Ser	Arg	Ser 80
	Ser	Ser	Leu	Gly	Asn 85	Lys	Ser	Pro	Gln	Leu 90	Ser	Gly	Asn	Leu	Ser 95	Gly
10	Gln	Ser	Ala	Ala 100	Ser	Val	Leu	His	Pro 105	Gln	Gln	Thr	Leu	His 110	Pro	Pro
15	Gly	Asn	Ile 115	Pro	Glu	Ser	Gly	Gln 120	Asn	Gln	Leu	Leu	Gln 125	Pro	Leu	Lys
		130					135					140		Ser		
20	145					150					155			Thr		160
٠					165					170				Gln	175	
25				180					185					190		Leu
30			195					200					205			Gly
		210	)				215					220	ı			Ala
35	225	5				230					235	i				Asn 240
					245	1				250	)				. 255	
40				260	)				265	j				270	)	Ala
45			279	5				280	)				289	5		Gln
		29	0				299	5			•	30	0			n Phe
50	30	5				1.Glr 310		s Se	r Ile	e Se:	r Ası 31		o Pr	o GT	y se	Asn 320
	Le:	u Ar	g Thi	r Thi	r											,
55																

- (2) INFORMATION FOR SEQ ID NO: 463:
  - (i) SEQUENCE CHARACTERISTICS:

60 (A) LENGTH: 133 amino acids

	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 463:
5	Ile Gln Asp Leu Gln Ser Arg Gln Lys His Glu Ile Glu Ser Leu Tyr 1 5 10 15
10	Thr Lys Leu Gly Lys Val Pro Pro Ala Val Ile Ile Pro Pro Ala Ala 20 25 30
	Pro Leu Ser Gly Arg Arg Arg Pro Thr Lys Ser Lys Gly Ser Lys 35 40 45
15	Ser Ser Arg Ser Ser Ser Leu Gly Asn Lys Ser Pro Gln Leu Ser Gly 50 60
	Asn Leu Ser Gly Gln Ser Ala Ala Ser Val Leu His Pro Gln Gln Thr 65 70 75 80
20	Leu His Pro Pro Gly Asn Ile Pro Glu Ser Gly Gln Asn Gln Leu Leu 85 90 95
25	Gln Pro Leu Lys Pro Ser Pro Ser Ser Asp Asn Leu Tyr Ser Ala Phe 100 105 110
	Thr Ser Asp Gly Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly Gln 115 120 125
30	Gly Thr Ser Ser Thr 130
35	(2) INFORMATION FOR SEQ ID NO: 464:
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 53 amino acids  (B) TYPE: amino acid
40	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 464:
	Thr Ser Asp Gly Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly Glr 1 5 10 15
45	Gly Thr Ser Ser Thr Asn Thr Val Gly Ala Thr Val Asn Ser Gln Ala 20 25 30
50	Ala Gln Ala Gln Pro Pro Ala Met Thr Ser Ser Arg Lys Gly Thr Pho 35 40 45
50	Thr Asp Asp Leu His 50
55	(2) INFORMATION FOR SEQ ID NO: 465:
	(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 48 amino acids

(B) TYPE: amino acid

	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 465:
5	Lys Gly His Met Asn Tyr Glu Gly Pro Gly Met Ala Arg Lys Phe Ser  1 5 10 15
	Ala Pro Gly Gln Leu Cys Ile Ser Met Thr Ser Asn Leu Gly Gly Ser 20 25 30
10	Ala Pro Ile Ser Ala Ala Ser Ala Thr Ser Leu Gly His Phe Thr Lys 35 40 45
15	
	(2) INFORMATION FOR SEQ ID NO: 466:
20	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 31 amino acids</li><li>(B) TYPE: amino acid</li></ul>
25	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 466:
	Gln Pro Leu Lys Pro Ser Pro Ser Ser Asp Asn Leu Tyr Ser Ala Phe  1 5 10 15
30	Thr Ser Asp Gly Ala Ile Ser Val Pro Ser Leu Ser Ala Pro Gly 20 25 30
35	(2) INFORMATION FOR SEQ ID NO: 467:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 57 amino acids
40	(A) LENGTH: 157 and no acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 467:
	Val Arg Val Ala Ala Ala Glu Ser Met Xaa Leu Leu Leu Glu Cys Ala 1 5 10 15
45	Xaa Val Arg Gly Pro Glu Tyr Leu Thr Gln Met Trp His Phe Met Cys 20 25 30
50	Asp Ala Leu Ile Lys Ala Ile Gly Thr Glu Pro Asp Ser Asp Val Leu 35 40 45  Ser Glu Ile Met His Ser Phe Ala Lys
55	50 55
"	

(2) INFORMATION FOR SEQ ID NO: 468:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 85 amino acids

60 (B) TYPE: amino acid

(D) TOPOLOGY: linear

			(xi)	SEQU	JENCI	E DES	CRIE	PTIO	N: SI	EQ II	ОИ C	468	3 :				
5	Met 1	Glu	Ile	Asn	Asn 5	Gln	Asn	Cys	Phe	Ile 10	Val	Ile	Asp	Leu	Val 15	Arg	
	Thr	Val	Met	Glu 20	Asn	Gly	Val	Glu	Gly 25	Leu	Leu	Ile	Phe	Gly 30	Ala	Phe	
10	Leu	Pro	Glu 35	Ser	Trp	Leu	Ile	Gly 40	Val	Arg	Cys	Ser	Ser 45	Glu	Pro	Pro	
15	Lys	Ala 50	Leu	Leu	Leu	Ile	Leu 55	Ala	His	Ser	Gln	Lys 60	Arg	Arg	Leu	Asp	
	Gly 65	Trp	Ser	Phe	Ile	Arg 70	His	Leu	Arg	Val	His 75	Tyr	Cys	Val	Ser	Leu 80	
20	Thr	Ile	His	Phe	Ser 85												
25	(2)	INF	ORMA	TION	FOR	SEQ	ID 1	<b>%</b> : 0	<b>4</b> 69:								
			(i)·	(	(A) L	CHAI ENGT YPE:	H: 2	0 am	ino		ls						
30			(xi)	(	(D) I	OPOL	OGY:	lin	ear	EQ I	D NO	: 46	9:				
	Gln 1		Lys	His	Ala 5		Glu	Val	Arg	Lys 10		Lys	Glu	Leu	. Lys 15	Glu	
35	Glu	. Ala	Ser	Arg 20													
40	(2)	INF	ORMA	TION	FOR	SEQ	ID:	NO:	470:				,	ı			
45			(i)		(A) I (B) :	CHA LENGI LYPE:	H: 9	92 ar ino a	mino acid	_	ds						
	• • • • • • • • • • • • • • • • • • • •	<b>~</b> 1				E DE								Dr.	, I.a.	ı Yaa	
50	1				. 5	•				10	)				15		
				20	)				25	•				30	)	ı Arg	
55			35	5				40	)				45	5		o Phe	
	Sei	Cy:		o Let	ı Ala	a Ser	55 55		e Val	l Pro	Gly	Glr 60		Cy:	s Va	l Thr	
60	Cys	s Pro	o Ph	e Pro	Sei	. Leu	Pro	Phe	e Glr	n Ası	p Lys	. His	ala	a Gl	ı Gl	u Val	

	55					70		•			75					80
_	Arg :	Lys .	Asn :	Lys :	Glu : 85	Leu :	Lys (	Glu (	Glu .	Ala 90	Ser .	Arg		•		
5																
•	(2)	250	FYAT	IC::	FOR :	SEQ	ID N	0: 4	71:							
10			i) S ki;	(3 (3 (3	E) LE E) TY C) TO	NGTH PE: POLO	H: 37 amir XGY:	7 ami no ac line	no a id ar	cid		471	.:			
15	Pro 1	Thr	Arg	Cys	Cys 5	Thr	Thr	Gln	Pro	Cys 10	Arg	Ser	Ser	Ala	Arg 15	Arg
20	Pro	≎ys	. طبي	Val 20	3ro ∶	Met	Val	Pro	Ser 25	Pro	Glu	Gly	Arg	Glu 30	Xaa	Gln
	Ero '		Cys 35	Pro	Ser									١		
25												·				
	(2)		FYAI													
30			(i)	() ()	A) LI B) T D) T	engt: YPE : OPOL	H: 3: ami: OGY:	63 ar no ac line	mino cid ear	aci		: 472	2:			
35	Met 1	Lys	yrg	Ser	Leu 5	Asin	Glu	Asn	Ser	Ala 10	Arg	Ser	Thr	Ala	Gly 15	Cys
40	Leu	220	Val	Pro 20	Leu	Phe	Asn	Gln	Lys 25	Lys	Arg	Asn	Arg	Gln 30	Pro	Leu
.••	Thr	Ser	Asn 35	Szo	Leu	Lys	Asp	Asp 40	Ser	Gly	Ile	Ser	Thr 45	Pro	Ser	Asp
45	Asn	20 تىدىت	Аsp	Phe	510	Pro	Leu 55	Pro	Thr	Asp	Trp	Ala 60	Trp	Glu	Ala	Val
	Asn 65	Pro	Glu	Χaa	Ala	Pro 70	Val	Met	Lys	Thr	Val 75	Asp	Thr	Gly	Gln	Ile 80
50	220	His	Ser	Val	Ser 85	Arg	Pro	Leu	Arg	Ser 90	Gln	Asp	Ser	Val	Phe 95	Asn
55	Ser	Ile	Gln	Ser 150	Asn	Thr	Gly	Arg	Ser 105	Gln	Gly	Gly	Trp	Ser 110	Tyr	Arg
	ನಿತ್ರಾ	Gly	Asn 115		Asn	Thr	Ser	Leu 120	Lys	Thr	Trp	Xaa	Lys 125	Asn	Asp	Phe
60	Lyš	Pro 130		Cys	Lys	Arg	Thr 135		Leu	Val	Ala	Asn 140	Asp	Gly	Lys	Asn

	Ser 145	Cys	Pro	Met	Ser	Ser 150	Gly	Ala	Gln	Gln	Gln 155	Lys	Gln	Leu	Arg	Thr 160	
5	Pro	Glu	Pro	Pro	Asn 165	Leu	Ser	Arg	Asn	Lys 170	Glu	Thr	Glu	Leu	Leu 175	Arg	
10	Gln	Thr	His	Ser 180	Ser	Lys	Ile	Ser	Gly 185	Cys	Thr	Met	Arg	Gly 190	Leu	Asp	
10	Lys	Asn	Ser 195	Ala	Leu	Gln	Thr	Leu 200	Lys	Pro	Asn	Phe	Gln 205	Gln	Asn	Gln	
15	Tyr	Lys 210	Xaa	Gln	Met	Leu	Asp 215	Asp	Ile	Pro	Glu	Asp 220	Asn	Thr	Leu	Lys	
	Glu 225	Thr	Ser	Leu	Tyr	Gln 230	Leu	Gln	Phẹ	Lys	Glu 235	Lys	Ala	Ser	Ser	Leu 240	
20	Arg	Ile	Ile	Ser	Ala 245		Ile	Glu	Ser	Met 250		Týr	Trp	Arg	Glu 255	His	
25	Ala	Gln	Lys	Thr 260		Leu	Leu	Phe	Glu 265	Val	Leu	Ala	Val	Leu 270		Ser	
	Ala	Val	Thr 275		Gly	Pro	Tyr	Тут 280		Lys	Thr	Phe	Leu 285		Arg	Asp	
30	Gly	Lys 290		Thr	Leu	Pro	Cys 295		Phe	Tyr	Glu	11e 300		Arg	Glu	Leu	
	Pro 305		Leu	ıIle	Arg	310		Val	. His	Arg	315		Gly	Asn	Tyr	320	
35	Glr	Lys	Lys	Asr	325		Gln	Cys	: Val	330		Arg	Pro	Ala	Ser 335	Val	
40	Ser	Glu	ı Glr	1 Lys 340		Phe	Glr	Ala	345		l Lys	: Ile	Ala	Asp 350		. Glu	
	Met	: Glr	тул 359		: Ile	e Asr	ı Val	. Met 360	Asr	Glu	ı Thi	•					
45	(2)	INI	FORM	ATIO	1 FOI	R SE(	Q ID	NO:	473:	1							
50					(A) (B) (D)	LENG TYPE TOPO	TH: : am LOGY	45 a ino : li	STIC: mino acid near ON:	aci		O: 4º	73:				
55		r Gli 1	n As	p Se	r Va		e Ası	n Se	r Ile	e Gl		r Ası	n Th	r Gl	y Ar	g Ser 5	:
	Gl	n Gl	y Gl	y Tr		г Ту	r Ar	g As	p Gly		n Ly	s As	n Th	r Se 3		u Lys	\$
60	Th	r Tr	рХа	a Ly	s As	n As	p Ph	e Ly	s Pr	o G1	n Cy	s Ly	s Ar	g			

45 35 5 (2) INFORMATION FOR SEQ ID NO: 474: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 36 amino acids (B) TYPE: amino acid 10 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 474: Asn Lys Glu Thr Glu Leu Leu Arg Gln Thr His Ser Ser Lys Ile Ser 5 15 Gly Cys Thr Met Arg Gly Leu Asp Lys Asn Ser Ala Leu Gln Thr Leu Lys Pro Asn Phe 20 35 (2) INFORMATION FOR SEQ ID NO: 475: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 49 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 475: 30 Ser Ser Leu Arg Ile Ile Ser Ala Val Ile Glu Ser Met Lys Tyr Trp Arg Glu His Ala Gln Lys Thr Val Leu Leu Phe Glu Val Leu Ala Val 35 Leu Asp Ser Ala Val Thr Pro Gly Pro Tyr Tyr Ser Lys Thr Phe Leu 40 40 Met 45 (2) INFORMATION FOR SEQ ID NO: 476: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 42 amino acids 50 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 476: Pro Arg Leu Ile Arg Gly Arg Val His Arg Cys Val Gly Asn Tyr Asp **55**. . 10 5 Gln Lys Lys Asn Ile Phe Gln Cys Val Ser Val Arg Pro Ala Ser Val

25

20

Ser Glu Gln Lys Thr Phe Gln Ala Phe Val

35	40
33	

5	(2)	INFC	RMAT	ION	FOR	SEQ	ID N	0: 4	77:							
10				(2 (1 (1	INCE A) LE B) TY O) TO JENCE	NGTH PE: POLC	I: 37 amir XGY:	0 an no ac line	mino cid ear	acid		477	:			
15	. 1				Pro 5					10					15	
	Ser	Pro	Leu	Asp 20	Pro	Glu	Val	Gly	Pro 25	Tyr	Cys	Asp	Thr	30	Thr	Met
20	Arg	Thr	Leu 35	Phe	Asn	Leu	Leu	Trp 40	Leu	Ala	Leu	Ala	Cys 45	Ser	Pro	Val
	His	Thr 50	Thr	Leu	Ser	Lys	Ser 55	Asp	Ala	Lys	Lys	Ala 60	Ala	Ser	Lys	Thr
25	Leu 65	Leu	Glu	Lys	Ser	Gln 70	Phe	Ser	Asp	Lys	Pro 75	Val	Gln	Asp	Arg	80
20	Leu	Val	Val	Thr	Asp 85	Leu	Lys	Ala	Glu	Ser 90	Val	Val	Leu	Glu	His 95	Arg
30	Ser	Tyr	Cys	Ser 100	Ala	Lys	Ala	Arg	Asp 105	Arg	His	Phe	Ala	Gly 110	Asp	Val
35	Leu	Gly	Tyr 115	Val	Thr	Pro	Trp	Asn 120		His	Gly	Tyr	Asp 125	Val	Thr	Lys
	Val	Phe 130		Ser	Lys	Phe	Thr 135		Ile	Ser	Pro	Val 140	Trp	Leu	Gln	Leu
40	Lys 145		Arg	Gly	Arg	Glu 150		Phe	Glu	Val	Thr 155		Leu	His	Asp	Val 160
45	Asp	Gln	Gly	Trp	Met 165		Ala	. Val	Arg	Lys 170		Ala	Lys	Gly	Leu 175	His
45	Ile	Val	. Pro	Arg 180								Tyr		Asp 190		Arg
50	Asr	Va]	. Leu 195		Ser	Glu	Asp	Glu 200		Glu	Glu	Leu	Ser 205		Thr	Val
	Val	Glr 210		. Ala	a Lys	Asn	Glr 215		: Phe	Asp	Gly	Phe 220		Val	. Glu	Val
55	Trg 225		ı Glr	ı Lev	ı Lev	Ser 230		ı Lys	arg	y Val	Gly 235		Ile	His	: Met	: Leu 240
60	Thi	r His	s Lev	ı Ala	a Glu 245		Let	ı His	3 Glr	1 Ala 250		, Leu	Leu	Ala	255	ı Leu

	Val	Ile	Pro	Pro 260	Ala	Ile	Thr	Pro	Gly 265	Thr	Asp	Gln	Leu	Gly 270	Met	Phe
5	Thr	His	Lys 275	Glu	Phe	Glu	Gln	Leu 280	Ala	Pro	Val	Leu	Asp 285	Gly	Phe	Ser
	Leu	Met 290	Thr	Tyr	Asp	Tyr	Ser 295	Thr	Ala	His	Gln	Pro 300	Gly	Pro	Asn	Ala
10	Pro 305	Leu	Ser	Trp	Val	Arg 310	Ala	Cys	Val	Gln	Val 315	Leu ;	Asp	Pro	Lys	Xaa 320
15	Lys	Trp	Arg	Thr	Lys 325	Ser	Ser	Trp	Gly	Ser 330	Thr	Ser	Met	Xaa	Trp 335	Thr
10	Xaa	Arg	Xaa	Pro 340	Xaa	Asp	Ala	Arg	Xaa 345	Pro	Val	Val	Gly	Xaa 350	Arg	Xaa
20	Ile	Gln	Xaa 355	Leu	Lys	Asp	His	Хаа 360	Pro	Arg	Met	Val	Leu 365	Asp	Ser	Lys
	Pro	Gln 370						·					٠			
25	(2)	INF	ORMA	TION	FOR	SEQ	ID:	NO:	478:				•			
20	,_,			SEQU	ENCE	СНА	RACT	ERIS	TICS		1_					
30				(	(A) I	.ENG1	H: 3	sy an	no	acıc	ເຣ					
			(xi)	(	(B) T	YPE:	ami : OGY	no a	cid near			): 47	8:			
35	Thr 1			(	(B) I (D) I	YPE: OPOI E DE	ami OGY: SCRI	no a : lir :PTIC	cid near N: S	EQ I	D NO			Asp	Thr 15	
	1		Ser	SEQ Pro	(B) T (D) T (UENC Leu 5	YPE: OPOI E DE	ami OGY: SCRI Pro	no a lir PTIC	ncid near ON: S	Gly 10	D NO	Tyr	Cys		15 . Cys	
35 40	1 Thr	Met	Ser Arg	SEQ Pro Thr 20	(B) T (D) T (UENC Leu 5	YPE: OPOI E DE Asp	ami OGY: SCRI Pro	ino a lir PTIC Glu	ncid near N: S Val	Gly 10	D NO	Tyr	Cys	Ala	15 . Cys	
	Thr	Met	Ser Arg His	SEQ Pro Thr 20	(B) T (D) T (UENC Leu 5 Leu Thr	TYPE: TOPOI E DE Asp Phe	ami OGY: SSCRI Pro Asn	no a lir PTIO Glu	cid near NN: S Val Leu 25	Gly 10	D NO	Tyr	Cys	Ala	15 . Cys	
40	Thr	Met	Ser Arg His 35	SECONDATION SEQUENCES	(B) 11 (D) 1	TYPE: COPOI  ASP  Phe  Leu  SEQ  CHECKER  CHECKER  CHECKER  CHECKER  COPOI  COPOR  COP	ami OGY: SCRI Pro Asn Ser	no a lir	acid near N: S Val Leu 25	Gly 10 Trp	Pro	Tyr	Cys	Ala	15 . Cys	
40	Thr	Met	Arg His 35	SECTION SEQUENTS	(B) 1 (D) 1 (D) 1 (D) 1 (D) 1 (E) 1 (E) 1 (E) 1 (D) 1	YPE: YPE: YPE: YPE: YPE: YPE: YPE: YPE:	ami OGY: SCRI Pro Asn Asn ID ARACT TH: : am	no a lir	acid near N: S Val Leu 25 479:	Gly 10 Trp	D NO	Tyr Ala	Cys	Ala	15 . Cys	
40 45	Thr Pro	Met	Arg His 35	SEQUENT SEQUENTS SEQU	(B) 1 (D) 1 (D) 1 (D) 1 (D) 1 (E) 1 (E) 1 (D) 1	YPE: YPE: YPE: YPE: YPE: YPE: YPE: YPE:	ami OGY: SCRI Pro Asn Asn ID ARACT H: : am LOGY	NO:  NO:  NO:  Inc. A control of the	acid near N: S Val Leu 25 479: strics mino acid near	Gly 10 Trp acid	D NO Pro Leu	Tyr Ala	Cys Leu	30 30	15	Ser
40 45 50	Thr Pro  (2)	Met Val INF	Arg His 35 CORMA (i) (xi	SEQUENT SEQUEN	(B) 11 (D) 11 (D) 17 (D	YPE: YPE: YPE: YPE: YPE: YPE: YPE: YPE:	ami OGY: SCRI Pro Asn Asn ID ARACT H: !: am LOGY	NO:  NO:  NO:  Inc. Ala.	acid hear N: S Val Leu 25	Gly 10 Gly 10 Trp acid	D NO Pro Leu	Tyr Ala	Cys Leu 79:	Ala 30	15 Cys	Ser

WO 98/54963

609

	35		40		45	
5	Val Phe Gly 50	Ser Lys Phe			·	
	(2) INFORMAT	TION FOR SEQ	ID NO: 480:			
10		(B) TYPE:	H: 52 amino a amino acid DGY: linear	acids	<b>4</b> 80:	
15	Arg Glu Met 1	Phe Glu Val	Thr Gly Leu	His Asp Va	al Asp Gln	Gly Trp
20	Met Arg Ala	Val Arg Lys 20	His Ala Lys 25	Gly Leu H	is Ile Val 30	Pro Arg
	Leu Leu Phe	Glu Asp Trp	Thr Tyr Asp 40	Asp Phe A	rg Asn Val 45	Leu Asp
25	Ser Glu Asp 50	Glu				
30		TION FOR SEQ				
35		(B) TYPE:	H: 56 amino amino acid  OGY: linear	acids	481:	
40	1	Gly Phe Val		10		15
	Lys Arg Val	l Gly Leu Ile 20	His Met Leu 25		Leu Ala Glu 30	
45	His Gln Ala	a Arg Leu Leu 5	Ala Leu Leu 40	Val Ile I	Pro Pro Ala 45	lle Thr
	Pro Gly Thi 50	r Asp Gln Leu	Gly Met 55			
50					:	
	(2) INFORM	ation for SEC	Į ĮD NO: 482:			
55		(B) TYPE	TH: 47 amino : amino acid LOGY: linear	acids	482:	

Asp Gly Phe Ser Leu Met Thr Tyr Asp Tyr Ser Thr Ala His Gln Pro

	. 1				5					10					15	
_	Gly	Pro	Asn	Ala 20	Pro	Leu	Ser	Trp	Val 25	Arg	Ala	Cys	Val	Gln 30	Val	Leu
5	Asp	Pro	Lys 35	Xaa	Lys	Tṛp	Arg	Thr 40	Lys	Ser	Ser	Trp	Gly 45	Ser	Thr	
10	(2)	INFO	ORMAT	rion	FOR	SEQ	ID N	IO: 4	183:							
15				(	A) Li B) T D) T	ENGTI YPE : OPOLA	H: 1 ami XGY:	52 a no a lin	mino cid ear	aci		: 48.	3:			
20	Glu 1	Arg	Gly	Val	Ser 5	Ile	Asn	Gln	Phe	Cys 10	Lys	Glu	Phe	Asn	Glu 15	Arg
	Thr	Lys ,	Asp	Ile 20	Lys	Glu	Gly	Ile	Pro 25	Leu	Pro	Thr	Lys	Ile 30	Leu	Val
25	Lys	Pro	Asp 35	Arg	Thr	Phe	Glu	Ile 40	Lys	Ile	Gly	Gln	Pro 45	Thr	Val	Ser
30	Tyr	Phe 50		Ľуs	Ala	Ala	Ala 55	Gly	Ile	Glu	Lys	Gly 60	Ala	Arg	Gln	Thr
. 50	Gly 65	Lys	Glu	Val	Ala	Gly 70	Leu	Val	Thr	Leu	Lys 75		Val	Tyr	Glu	Ile 80
35	Ala	Arg	Ile	Lys	Ala 85	Gln	Asp	Glu	Ala	Phe 90		Leu	Gln	Asp	Val 95	Pro
	Leu	Ser	Ser	Val 100	Val	Arg	Ser	[ Ile	11e 105		Ser	Ala	Arg	Ser 110		Gly
40	Ile	Arg	Val 115		Lys	Asp	Leu	Ser 120		Glu	Glu	Leu	Ala 125		Phe	Gln
45		130	)	Ala Glu			135	•		Gln	Lys	140		Asp	Leu	ı Ala
	145					150										
50	(2)	IN	ORM	VTION	FOF	SEC	) ID	NO:	484:							_
55		-			(A) : (B) ' (D) '	LENG TYPE TOPO	TH: : am LOGY	270 ino : li	amin acid near	o ac		D: 48	34:			
60	Ala		1 Ty:	r Thi	Ty:		s Gl	ı Ly:	s Lys	S Lys 10		7h1	Ala	a Ala	a Se:	r Gly

	Tyr	Gly	Thr	Gln 20	Asn	Ile	Arg	Leu	Ser 25	Arg	Asp	Ala	Val	Lys 30	Asp	Phe
5	Asp	Суз	Cys 35	Cys	Leu	Ser	Leu	Gln 40	Pro	Cys	His	Asp	Pro 45	Val	Val	Thr
10	Pro	Asp 50		Tyr	Leu	Tyr	Glu 55	Arg	Glu	Ala	Ile	Leu 60	Glu	Tyr	Ile	Leu
10	His 65	Gln	Lys	Lys	Glu	Ile 70	Ala	Arg	Gln	Met	Lys 75	Ala	Tyr	Glu	Lys	Gln 80
15	Arg	Gly	Thr	Arg	Arg 85	Glu	Glu	Gln	Lys	Glu 90	Leu	Gln	Arg	Ala	Ala 95	Ser
	Gln	Asp	His	Val 100	Arg	Gly	Phe	Leu	Glu 105	Lys	Glu	Ser	Ala	Ile 110	Val	Ser
20	Arg	Pro	Leu 115		Pro	Phe	Thr	Ala 120	Lys	Ala	Leu	Ser	Gly 125	Thr	Ser	Pro
25	Asp	Asp 130		Gln	Pro	Gly	Pro 135	Ser	Val	Gly	Pro	Pro 140		Lys	Asp	Lys
23	Asp 145		Val	Leu	Pro	Ser 150		Trp	Ile	Pro	Ser 155		Thr	Pro	Glu	Ala 160
30	Lys	Ala	Thr	Lys	Leu 165		Lys	Pro	Ser	170		Val	Thr	Cys	Pro 175	Met
				180	)				185	•				190		Thr
35			199	5			•	200	)				209	5		Glu
40		210	)				21	5				220	)			Pro
	225	5	•			23	0				239	5 .				240
45					24	5				25	0				25!	b rys
	Let	ı Th	r As	p Ar		p Il	e Il	e Va	1 Le: 26:		n Ar	g Gly	y Gl	y Thi 270	r 0	
50							•									
	(2	) IN	FORM	ATIO	n fo	R SE	QII	NO:	485	:						
55					(A) (B) (D)	TYPE TOPO	FTH: E: au OLOG	TERI 54 a mino Y: li	mino ació near	aci i						
			(xi	L) SE	QUE	ICE I	DESCI	RIPTI	ON:	SEQ	ID 1	10:4	85 :			•

Tyr Leu Tyr Glu Arg Glu Ala Ile Leu Glu Tyr Ile Leu His Gln Lys

	1				5					10					15	
5	Lys	Glu	Ile	Ala 20	Arg	Gln	Met	Lys	Ala 25	Tyr	Glu	Lys	Gln	Arg 30	Gly	Thr
5	Arg	Arg	Glu 35	Glu	Gln	Lys	Glu	Leu 40	Gln	Arg	Ala	Ala	Ser 45	Gln	Asp	His
10	Val	Arg 50	Gly	Phe	Leu	Glu										
15	(2)	INF														
			(i)	(	A) L B) T	ENGT YPE:	H: 6	4 an	ino cid		ls					
20			(xì)	SEQ				PTIC		EQ I	D NC	: 48	6:			
	Phe 1	Thr	Ala	Lys	Ala 5		Ser	Gly	Thr	Ser 10		Asp	Asp	Val	Gln 15	Pro
25	Gly	Pro	Ser	Val 20		Pro	Pro	Ser	Lys 25		Lys	Asp	Lys	Val 30		Pro
30	Ser	Phe	Trp 35		Pro	Ser	Leu	Thr 40		Glu	a Ala	Lys	Ala 45		Lys	Leu
50	Glu	Lys 50		Ser	Arg	Thr	Va]	_	Cys	Pro	Met	Ser 60		Lys	Pro	Leu
35	,	,					-									•
40	(2)	) INI											ě			
•			(1)	SEQ	(A) (B)	LENG TYPE	TH: : an	56 a nino	mino acid	aci	ds Î					
45			(xi	) SE				: li IPTI			ID N	0: 4	87:			
		1 Hi: 1	s Ph	e Th		o Le	u As	p Se	r Se	r Va 1		p Ar	g Va	l Gl	y Let 19	ı Ile 5
50	Th	r Ar	g Se	r Gl		g Ty	r Va	l Cy	s Al		l Th	r Ar	g As	p Se		ı Ser
55	As	n Al		r Pr 5	о Су	s Al	a Va		u Ar 0	g Pr	o Se	r Gl		a Va 5	l Va	l Thr
JJ	Le	eu G1 5	u Cy 0	s Va	l Gl	u Ly		eu Il i5	.e ·						•	

	(2)	INFC	RMAT	ION	FOR	SEQ	ID N	io: 4	88:							
5				() (1 (1	A) Li B) T D) T	ENGTI YPE : OPOLA	H: 50 amii OGY:	67 ar no ac line		acio		: 488	3:			
10	Met 1	Asp	Thr	Ser	Glu 5	Asn	Arg	Pro	Glu	Asn 10	Asp	Val	Pro	Glu	Pro 15	Pro
	Met	Pro	Ile	Ala 20	Asp	Gln	Val	Ser	Asn 25	Asp	Asp	Arg	Pro	Glu 30	Gly	Ser
15	Val	Glu	Asp 35	Glu	Glu	Lys	Lys	Glu 40	Ser	Ser	Leu	Pro	Lys 45	Ser	Phe	Lys
20	Arg	Lys 50	Ile	Ser	Val	Val	Ser 55	Ala	Thr	Lys	Gly	Val 60	Pro	Ala	Gly	Asn
20	Ser 65	Asp	Thr	Glu	Gly	Gly 70	Gln	Pro	Gly	Arg	Lys 75	Arg	Arg	Trp	Gly	Ala 80
25	Ser	Thr	Ala	Thr	Thr 85	Gln	Lys	Lys	Pro	Ser 90	Ile	Ser	Ile	Thr	Thr 95	Glu
	Ser	Leu	Lys	Ser 100		Ile	Pro	Asp	Ile 105	Lys	Pro	Leu	Ala	Gly 110	Gln	Glu
30	Ala	Val	Val 115	Asp	Leu	His	Ala	Asp 120	Asp	Ser	Arg	Ile	Ser 125	Glu	Asp	Glu
25	Thr	Glu 130		Asn	Gly	Asp	Asp 135		Thr	His	Asp	Lys 140	Gly	Leu	Lys	Ile
35	Cys 145		Thr	Val	Thr	Gln 150		Val	Pro	Ala	Glu 155		Gln	Glu	Asn	Gly 160
40	Gln	Arg	Glu	Glu	Glu 165		Glu	Glu	Lys	Glu 170		Glu	Ala	Glu	Pro 175	
	Val	Pro	Pro	Gln 180		Ser	Val	Glu	Vàl 185	Ala	Leu	Pro	Pro	Pro 190		Glu
45	His	Glu	Val 195		: Lys	Val	Thr	Leu 200		Asp	Thr	Leu	Thr 205		Arg	Ser
50	<u>I</u> le	Ser 210		Glr	Lys	: Ser	Gly 215		. Ser	Ile	. Thr	220		Asp	Pro	Val
50	Arg 225		: Ala	Glr	ı Val	230		Pro	Pro	Arg	Gly 235		Ile	Ser	Asn	1le 240
55	Val	. His	s Ile	e Sei	245		ı Val	l Arg	, Pro	250		Lev	ı Gly	Gln	Leu 255	Lys
	Glu	ı Leı	ı Leu	Gly 260		y Thi	c Gly	y Thi	Leu 265		l Glu	ı Glu	ı Ala	Phe 270		lle

Asp Lys Ile Lys Ser His Cys Phe Val Thr Tyr Ser Thr Val Glu Glu

			275					280					285			
~	Ala	Val 290		Thr	Arg	Thr .	Ala 295	Leu	His	Gly	Val	Lys 300	Trp	Pro	Gln	Ser
5	Asn 305	Pro	Lys	Phe	Leu	Cys 310	Ala	Asp	Tyr	Ala	Glu 315	Gln	Asp	Glu	Leu	Asp 320
10	Tyr	His	Arg	Gly	Leu 325	Leu	Val	Asp	Arg	Pro 330	Ser	Glu	Thr	Lys	Thr 335	Glu
	Glu	Gln	Gly	Ile 340	Pro	Arg	Pro	Leu	His 345	Pro	Pro	Pro	Pro	Pro 350	Pro	Val
15	Gln	Pro	Pro 355		His	Pro	Arg	Ala 360	Glu	Gln	Arg	Glu	Gln 365	Glu	Arg	Ala
20	Val	Arg 370		Gln	Trp	Ala	Glu 375	Arg	Glu	Arg	Glu	Met 380	Glu	Arg	Arg	Glu
20	Arg 385		: Arg	Ser	Glu	Arg 390	Glu	Trp	Asp	Arg	Asp 395	Lys	Val	Arg	Glu	Gly 400
25					405					410	1				415	
				420					425					430		Glu
30			435	5				440	)				445			Ala
35		45	0				455	5				460	)			Lys
	465	5				470					475	5				1 Lys 480
40					489	5				49	0	•			495	
				50	0				50:	5				510	)	r Arg
45			51	.5				52	0		•		525	5		y Asp
50	Ar		sp Ar 30	g As	p Ar	g Glı	53		p Ar	g Ģl	u Ar	g G1; 54	y Arg	g Gl	ı Ar	g Asr
	Ar 54		rg As	sp Th	r Ly	s . Ar:		s Se	r Ar	g Se	r Ar 55	g Se 5	r Ar	g Se	r Th	r Pro 560
55	Va	ıl A	rg As	sp Ax	g Gl 56	y G1; 5	y Ar	g								

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 489:
5
     Gly Cys Asp Ser Cys Pro Pro His Leu Pro Arg Glu Ala Phe Ala Gln
     Asp Thr Gln Ala Glu Gly Glu Cys Ser Ser Arg Ala Glu Arg Ala Asp
10
     Met Cys Pro Asp Ala Pro Pro Ser Gln Glu Val Pro Glu Gly Pro Gly
                                  40
15
     Ala Ala Pro
          50
20
      (2) INFORMATION FOR SEQ ID NO: 490:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 50 amino acids
25
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 490:
      Pro Gln Leu Pro Ser Cys Gly Arg Pro Trp Pro Gly Thr Ala Ser Val
30
      Phe Gln Ser His Thr Gln Gly Pro Arg Glu Asp Pro Asp Pro Cys Arg
      Ala Gln Gly Ser Ala Gly Thr His Cys Pro Ile Ser Leu Ser Pro Pro
35
                                  40
      Arg Gln
           50
40
       (2) INFORMATION FOR SEQ ID NO: 491:
45
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 42 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 491:
50
      Pro Gly Phe Arg Gly Pro Ser Gly Ser Leu Gly Cys Ser Phe Phe Pro
      Arg Ser Leu Gly Arg Val Leu Pro Pro Gly Cys Gln Arg Pro Gly Ala
55
                                    25
      His Ala Asp Ser Ser Pro Pro Pro Thr Pro
               35
```

	(2) INFORMATION FOR SEQ ID NO: 492:  (i) SEQUENCE CHARACTERISTICS:															
5				(, ()	A) Li B) T D) T	ENGTI YPE : OPOLA	H: 84 amii OGY:	4 am: no ac line	ino a cid ear	acids		: 492	2:			
10	Glu 1	Asp	Leu	Lys	Lys 5	Pro	Asp	Pro	Ala	Ser 10	Leu	Arg	Ala	Ala	Ser 15	Cys
15	Gly	Glu	Gly	Lys 20	Lys	Arg	Lys	Ala	Cys 25	Lys	Asn	Cys	Thr	<u>.C</u> ys 30	Gly	Leu
13	Ala	Glu	Glu 35	Leu	Glu	Lys	Glu	Lys 40	Ser	Arg	Glu	Gln	Met 45	Ser	Ser	Gln
20	Pro	Lys 50		Ala	Cys	Gly	Asn 55	Cys	Tyr	Leu	Gly	Asp 60	Ala	Phe	Arg	Cys
	Ala 65		Cys	Pro	туг	Leu 70		Met	Pro	Ala	Phe 75	Lys	Pro	Gly	Glu	Lys 80
25	Val	Leu	. Leu	. Ser						•						
30	(2)	INI		ATION SEQU												
35					(A) 1 (B) 1 (D) 1	ENG TYPE TOPOI	rh: 9 : am: Logy	90 ar ino a : lir	mino acid near	ació		): <b>4</b> 9	93:			
40	Glu		p Lei	u Lys	Lys		Asr	Pro	Ala	Ser 10		Arg	, Ala	Ala	Ser 15	Cys
40	Gl	y Gl	u Gl	y Ly: 20		Arg	J Lys	s Ala	Cys 25		: Asn	Cys	Thr	: Cys	Gly	/ Leu
45	Ala	a Gl	u Gl 3	_	u Glu	ı Lys	s Glu	1 Lys 40		Arg					Sei	Gln
	Pr	_	s Se O	r Al	a Cy:	s Gly	y Ası 5		з Туг	Leu	ı Gly	Ası 60		Phe	e Arg	g Cys
50	A1 6		r Cy	s Pr	о Ту:	r Lei		y Me	t Pro	Ala	a Phe 79		s Pro	Gly	/ Gl	80 L Lys
55	Va	l Le	eu Le	eu Se	r As		r As	n Le	u His	ASI 90		٠				

(2) INFORMATION FOR SEQ ID NO: 494:

(i) SEQUENCE CHARACTERISTICS:

```
(A) LENGTH: 34 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 494:
5
     Cys Gly Asn Cys Tyr Leu Gly Asp Ala Phe Arg Cys Ala Ser Cys Pro
     Tyr Leu Gly Met Pro Ala Phe Lys Pro Gly Glu Lys Val Leu Leu Ser
                                       25
10
                   20
      Asp Ser
15
      (2) INFORMATION FOR SEQ ID NO: 495:
            (i) SEQUENCE CHARACTERISTICS:
20
                     (A) LENGTH: 25 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 495:
      Ser Cys Gly Glu Gly Lys Lys Arg Lys Ala Cys Lys Asn Cys Thr Cys
25
                        5
        1
      Gly Leu Ala Glu Glu Leu Glu Lys Glu
                   20
30
       (2) INFORMATION FOR SEQ ID NO: 496:
35
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 21 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 496:
40
      Ser Gln Pro Lys Ser Ala Cys Gly Asn Cys Tyr Leu Gly Asp Ala Phe
                        5
      Arg Cys Ala Ser Cys
45
                    20
       (2) INFORMATION FOR SEQ ID NO: 497:
 50
              (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 17 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 497:
 55
       Arg Glu Ala Gly Gln Asn Ser Glu Arg Gln Tyr Val Ser Leu Ser Arg
 60
```

5	(2) INFORMATION FOR SEQ ID NO: 498:	
10	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 90 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 498:</li> </ul>	
15	Glu Ser Ser Gly Gln Ala Arg Thr Leu Ala Asp Pro Gly Pro Gly Trp 1 5 10 15	
15	Pro Arg Gln Gln Gly Met Cys Phe Gly Ser Leu Thr Gly Leu Ser Thr 20 25 30	
20	Thr Pro His Gly Phe Leu Thr Val Ser Ala Glu Ala Asp Pro Arg Leu 35 40 45	
	Ile Glu Ser Leu Ser Gln Met Leu Ser Met Gly Phe Ser Asp Glu Gly 50 55 60	
25	Gly Trp Leu Thr Arg Leu Leu Gln Thr Lys Asn Tyr Asp Ile Gly Ala 65 70 75 80	
30	Ala Leu Asp Thr Ile Gln Tyr Ser Lys His 85 90	
	(2) INFORMATION FOR SEQ ID NO: 499:	
35	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 159 amino acids  (B) TYPE: amino acid	
40	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 499:  Gln Glu Gly Ser Glu Pro Val Leu Leu Glu Gly Glu Cys Leu Val Val	
	1 5 10 15	
45	Cys Glu Pro Gly Arg Ala Ala Ala Gly Gly Pro Gly Gly Ala Ala Leu 20 25 30	L
	Gly Glu Ala Pro Pro Gly Arg Val Ala Phe Xaa Ala Val Arg Ser His 35 40 45	}
50	His His Glu Pro Ala Gly Glu Thr Gly Asn Gly Thr Ser Gly Ala Ile 50 55 60	<u> </u>
55	Tyr Phe Asp Gln Val Leu Val Asn Glu Gly Gly Phe Asp Arg Ala 65 70 75 86	æ O
	Ser Gly Ser Phe Val Ala Pro Val Arg Gly Val Tyr Ser Phe Arg Pho 85 90 95	9
60	His Val Val Lys Val Tyr Asn Arg Gln Thr Val Gln Val Ser Leu Me	t

	Leu Asn Thr Trp Pro Val Ile Ser Ala Phe Ala Asn Asp Pro Asp Val 115 120 125
5	Thr Arg Glu Ala Ala Thr Ser Ser Val Leu Leu Pro Leu Asp Pro Gly 130 135 140
10	Asp Arg Val Ser Leu Arg Leu Arg Gly Xaa Ser Thr Gly Trp 145 150 155
	(2) INFORMATION FOR SEQ ID NO: 500:
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 32 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 500:
	Pro Arg Ser Arg Pro Ala Leu Arg Pro Gly Arg Gln Arg Pro Pro Ser 1 5 10 15
25	His Ser Ala Thr Ser Gly Val Leu Arg Pro Arg Lys Lys Pro Asp Pro 20 25 30
30	
	(2) INFORMATION FOR SEQ ID NO: 501:
35	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 31 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 501:</li> </ul>
40	Met Thr Leu Ile Thr Pro Ser Xaa Lys Leu Thr Phe Xaa Lys Gly Asn
	1 5 10 15
45	Lys Ser Trp Ser Ser Arg Ala Cys Ser Ser Thr Leu Val Asp Pro 20 25 30
	(2) INFORMATION FOR SEQ ID NO: 502:
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 51 amino acids  (B) TYPE: amino acid
م م	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 502:
55	Gly His Pro Ser Pro Ala Leu Ser Ile Ala Pro Ser Asp Gly Ser Gln 1 5 10 15
60	Leu Pro Cys Asp Glu Val Pro Tyr Gly Glu Ala His Val Thr Arg Tyr 20 25 30

	Cys Lys Lys Pro Leu Thr Asn Ser His Leu Glu Thr Glu Ala Gln Ser 35 40 45	
5	Ser Ser Leu 50	
		-
10	(2) INFORMATION FOR SEQ ID NO: 503:	
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 263 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 503:	
20	GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC	60
	GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT	120
25	CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC	180
23	TCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	240
	AAGTATTAAA AGTAGCTITG TAA	263
30		
	(2) INFORMATION FOR SEQ ID NO: 504:	٠
35	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 263 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 504:	
	GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC	60
45	GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT	120
	CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC	180
50	TCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	240
50	AAGTATTAAA AGTAGCTTTG TAA	263
55	(2) INFORMATION FOR SEQ ID NO: 505:	
	(i) SEQUENCE CHARACTERISTICS:	
60	(A) LENGTH: 263 base pairs (B) TYPE: nucleic acid	

	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 505:	
5	GCTTCGTGTC CAACCCTCTT GCCCTTCGCC TGTGTGCCTG GAGCCAGTCC CACCACGCTC	60
	GCGTTTCCTC CTGTAGTGCT CACAGGTCCC AGCACCGATG GCATTCCCTT TGCCCTGAGT	120
10	CTGCAGCGGG TCCCTTTTGT GCTTCCTTCC CCTCAGGTAG CCTCTCTCCC CCTGGGCCAC	180
	TCCCGGGGGT GAGGGGGTTA CCCCTTCCCA GTGTTTTTTA TTCCTGTGGG GCTCACCCCA	240
1.5	AAGTATTAAA AGTAGCTTTG TAA	263
15		
20	(2) INFORMATION FOR SEQ ID NO: 506:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 160 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double	
25	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 506:	
30	TGGCTCACTG TCTTACAATC ACTGCTGTGG AATCATGATA CCACTTTTAG CTCTTTGCAT	60
	CTTCCTTCAG TGTATTTTTG TTTTTCAAGA GGAAGTAGAT TTTAACTGGA CAACTTTGAG	120
	TACTGACATC ATTGATAAAT AAACTGGCTT GTGGTTTCAA	160
35		
	(2) INFORMATION FOR SEQ ID NO: 507:	
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 292 amino acids  (B) TYPE: amino acid	
45	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 507:	
	Leu Asp Glu Leu Met Ala His Leu Thr Glu Met Gln Ala Lys Val Ala  1 5 10 15	•
50	Val Arg Ala Asp Ala Gly Lys Lys His Leu Pro Asp Lys Gln Asp His 20 25 30	
	Lys Ala Ser Leu Asp Ser Met Leu Gly Gly Leu Glu Glu Leu Gln 35 40 45	
55	Asp Leu Gly Ile Ala Thr Val Pro Lys Gly His Cys Ala Ser Cys Gln 50 55 60	
60	Lys Pro Ile Ala Gly Lys Val Ile His Ala Leu Gly Gln Ser Trp His 65 70 75 80	

	Pro	Glu	His	Phe	Val 85	Cys	Thr	His	Cys	Lys 90	Glu	Glu	Ile	Gly	Ser 95	Ser
5	Pro	Phe	Phe	Glu 100	Arg	Ser	Gly	Leu	Xaa 105	Tyr	Cys	Pro	Asn	Asp 110	Tyr	His
	Gln	Leu	Phe 115	Ser	Pro	Arg	Cys	Ala 120	Tyr	Cys	Ala	Ala	Pro 125	Ile	Leu	Asp
10	Lys	Val 130	Leu	Thr	Ala	Met	Asn 135	Gln	Thr	Ттр	His	Pro 140	Glu	His	Phe	Phe
15	Cys 145	Ser	His	Cys	Gly	Glu 150	Val	Phe	Gly	Ala	Glu 155	Gly	Phe	His	Glu	Lys 160
13	Asp	Lys	Lys	Pro	Туг 165	Cys	Arg	Lys	Asp	Phe 170		Ala	Met	Phe	Ser 175	Pro
20	Lys	Cys	Gly	Gly 180	Cys	Asn	Arg	Pro	Val 185	Leu	Glu	Asn	Tyr	Leu 190	Ser	Ala
	Met	Asp	Thr 195	Val	Trp	His	Pro	Glu 200	Cys	Phe	Val	Cys	Gly 205		Cys	Phe
25	Thr	Ser 210		Ser	Thr	Gly	Ser 215		Phe	Glu	Leu	Asp 220	Gly	Arg	Pro	Phe
30	Cys 225		Leu	His	Tyr	His 230		Arg	Arg	Gly	Thr 235		Cys	His	Gly	Cys 240
50	Gly	Gln	Pro	Ile	Thr 245		Arg	Cys	Ile	Ser 250		. Met	Gly	Tyr	Lys 255	Phe
35	His	Pro	Glu	His 260		. Val	. Cys	ala	Phe 265		: Leu	Thr	Gln	270	Ser	Lys
	Gly	Ile	275		g Glu	Glr	a Asr	Asp 2,80		Thi	туг	Cys	Glr 285		Cys	Phe
40	Asn	Lys 290	Leu )	ı Ph∈	•											
45			70574	. MT ()	ı no			NO.	509							
43	(2)	INI						NO:								
•			(1)	SEQ	(A)	LENG	TH:	TERI 43 a	mino	aci	.ds					
50			(xi	) SE	(D)	TOPO	LOGY	ino : li IPTI	near		ID N	0: 5	08:			
<i>e e</i>		s Al	a Se	r Le		p Se 5	r Me	t Le	u Gl		y Le	u Gl	ı Gl	n Gli	ı Le	u Gln 5
55	Asj	p Le	u Gl	y Il		a Th	r Va	l Pr	o Ly 2		y Hi	s Cy	s Al	a Se	r Cy O	s Gln
60	Ly	s Pr		e Al 5	a Gl	у Lу	s Va	1 I1 4		s Al	a Le	u				

_	(2) INFORMATION FOR SEQ ID NO: 509:
5	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 50 amino acids</li><li>(B) TYPE: amino acid</li></ul>
	(D) TOPOLOGY: linear
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 509:
	Cys Pro Asn Asp Tyr His Gln Leu Phe Ser Pro Arg Cys Ala Tyr Cys  1 5 10 15
15	Ala Ala Pro Ile Leu Asp Lys Val Leu Thr Ala Met Asn Gln Thr Trp 20 25 30
20	His Pro Glu His Phe Phe Cys Ser His Cys Gly Glu Val Phe Gly Ala 35 40 45
	Glu Gly 50
25	(2) INFORMATION FOR SEQ ID NO: 510:
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 67 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 510:
35	Asp Lys Lys Pro Tyr Cys Arg Lys Asp Phe Leu Ala Met Phe Ser Pro 1 5 10 15
	Lys Cys Gly Gly Cys Asn Arg Pro Val Leu Glu Asn Tyr Leu Ser Ala 20 25 30
40	Met Asp Thr Val Trp His Pro Glu Cys Phe Val Cys Gly Asp Cys Phe 35 40 45
45	Thr Ser Phe Ser Thr Gly Ser Phe Phe Glu Leu Asp Gly Arg Pro Phe 50 55 60
40	Cys Glu Leu 65
50	·
	(2) INFORMATION FOR SEQ ID NO: 511:
55	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 46 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 511:</li> </ul>
	Cys Gly Gln Pro Ile Thr Gly Arg Cys Ile Ser Ala Met Gly Tyr Lys
60	1 5 10 15

60

	Phe His	Pro (	Glu F 20	His 1	Phe \	Val (	Cys 1	Ala 25	Phe (	Cys 1	Leu '	Thr	Gln 30	Leu :	Ser
5	Lys Gly	Ile 35	Phe A	Arg (	Glu (	Gln i	Asn A	Asp	Lys '	Thr '	Tyr	Cys 45	Gln		
10	(2) INF							•							
15		(i) S	( <i>P</i> (E (I	) TY 3) TY	NGTH PE: POLC	I: 45 amir XGY:	52 am no ac line	nino :id ear	acio		512	2:			
20	Met Gly			5					10					15	
	Tyr His	val	Leu 20	Arg	Val	Gln	Glu	Asn 25	Ser	Pro	GIÀ	His	Arg 30	Ala	GIY
25	Leu Glu	ı Pro 35	Phe	Phe	Asp	Phe	Ile 40	Val	Ser	Ile	Asn	Gly 45	Ser	Arg	Leu
	Asn Ly		Asn	Asp	Thr	Leu 55	Lys	Asp	Leu	Leu	Lys 60	Xaa	Asn	Val	Glu
30	Lys Pr	o Val	Lys	Met	Leu 70	Ile	Tyr	Ser	Ser	Lys 75	Thr	Leu	Glu	Leu	Arg 80
25	Glu Th	r Ser	Val	Thr 85	Pro	Ser	Asn	Leu	Ттр 90	Gly	Gly	Gln	Gly	<b>Leu</b> 95	Leu
35	Gly Va	l Ser	Ile 100	Arg	Phe	Cys	Ser	Phe 105		Gly	Ala	Asn	Glu 110	Asn	Val
40	Trp Hi	s Val 115		Glu	Val	Glu	Ser 120		Ser	Pro	Ala	Ala 125		Ala	Gly
	Leu Ar 13	0				135		٠			140	1			
45	Glu Se 145	er Glu	a Asp	Leu	Phe 150		Leu	Ile	Glu	155	His	Glu	Ala	Lys	160
	Leu Ly	s Lev	1 Тух	Val 165		Asn	Thr	Asp	Thr 170	Asp	) Asr	ı Cys	Arg	g Glu 175	val
50	Ile I	le Thi	180		. Ser	: Ala	a Trp	Gly 185		y Glu	Gly	/ Sei	. Let 190	ı Gly	y Cys
55	Gly I	le Gly 199		Gly	туг	Leu	1 His 200		g Ile	e Pro	Th:	205	g Pro	o Phe	e Glu
	Glu G	ly Ly: 10	s Lys	s Ile	e Sei	Let 21		Gly	y Gli	n Met	220	a Gly	y Th	r Pro	o Ile

Thr Pro Leu Lys Asp Gly Phe Thr Glu Val Gln Leu Ser Ser Val Asn

	225					230					235					240
_	Pro	Pro	Ser	Leu	Ser 245	Pro	Pro	Gly	Thr	Thr 250	Gly	Ile	Glu	Gln	Ser 255	Leu
5	Thr	Gly	Leu	Ser 260	Ile	Ser	Ser	Thr	Pro 265	Pro	Ala	Val	Ser	Ser 270	Val	Leu
10	Ser	Thr	Gly 275	Val	Pro	Thr	Val	Pro 280	Leu	Leu	Pro		Gln 285	Val	Asn	Gln
	Ser	Leu 290	Thr	Ser	Val	Pro	Pro 295	Met	Asn	Pro	Ala	Thr 300	Thr	Leu	Pro	Gly
15	Leu 305	Met	Pro	Leu	Pro	Ala 310	Gly	Leu	Pro	Asn	Leu 315	Pro	Asn	Leu	Asn	Leu 320
20	Asn	Leu	Pro	Ala	Pro 325	His	Ile	Met	Pro	Gly 330	Val	Gly	Leu	Pro	Glu 335	Leu
	Val	Asn	Pro	Gly 340	Leu	Pro	Pro	Leu	Pro 345	Ser	Met	Pro	Pro	Arg 350	Asni	Leu
25	Pro	Gly	Ile 355	Ala	Pro	Leu	Pro	Leu 360	Pro	Ser	Glu	Phe	Leu 365	Pro	Ser	Phe
	Pro	Leu 370	Val	Pro	Glu	Ser	Ser 375	Ser	Ala	Ala	Ser	Ser 380	Gly	Glu	Leu	Leu
30	Ser 385	Ser	Leu	Pro	Pro	Thr 390	Ser	Asn	Ala	Pro	Ser 395	Asp	Pro	Ala	Thr	Thr 400
35	Thr	Ala	Lys	Ala	Asp 405	Ala	Ala	Ser	Ser	Leu 410	Thr	Val	Asp	Val	Thr 415	Pro
	Pro	Thr	Ala	Lys 420		Pro	Thr	Thr	Val 425	Glu	Asp	Arg	Val	Gly 430	Asp	Ser
40	Thr	Pro	Val 435	Ser	Glu	Lys	Pro	Val 440		Ala	Ala	Val	Asp 445	Ala	Asn	Ala
	Ser	Glu 450	Ser	Pro												
45	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	513:							
50				SEQU	ENCE		RACT	ERIS	TICS		ds					
			(xi)	(	(B) 1 (D) 1	YPE:	ami OGY:	ino a	acid near			): 51	.3 :			•
55	Ser 1		Glu	Ile	Pro		Gly	Gly	Thr	Glu 10		Туг	His	Val	Leu 15	Arg
60	Val	Gln	Glu	Asn 20		Pro	Gly	' His	Arg 25		Gly	Leu	Glu	Pro 30		Phe.

	Asp	Phe	Ile 35	Val	Ser	Ile	Asn	Gly 40	Ser	Arg	Leu	Asn	Lys 45	Asp	Asn	Asp
5	Thr	Leu 50	Lys	Asp	Leu	Leu	Lys 55	Xaa	Asn	Val	Glu	Lys 60	Pro	Val	Lys	Met
	Leu 65	Ile	Tyr	Ser	Ser	Lys 70	Thr	Leu	Glu	Leu	Arg 75	Glu	Thr	Ser	Val	Thr 80
10	Pro	Ser	Asn	Leu	Trp 85	Gly	Gly	Gln	Gly	Leu 90	Leu	Gly	Val	Ser	Ile 95	Arg
15	Phe	Cys	Ser	Phe 100	Asp	Gly	Ala	Asn	Glu 105	Asn	Val	Trp	His			
	(2)	INFO	ORMA!	rion	FOR	SEQ	ID I	NO: 5	514:							
20			(i)	(	A) L B) T	CHAI ENGT YPE: OPOL	H: 1 ami	45 a no a	mino cid		ds					
25				SEQ						_						
	Glu 1	Ser	Asn	Ser	Pro 5	Ala	Ala	Leu	Ala	Gly 10	Leu	Arg	Pro	His	Ser 15	Asp
30	Tyr	Ile	Ile	Gly 20	Ala	Asp	Thr	Val	Met 25	Asn	Glu	Ser	Glu	Asp 30	Leu	Phe
	Ser	Leu	Ile 35	Glu	Thr	His	Glu	Ala 40	Lys	Pro	Leu	Lys	Leu 45	Tyr	Val	Tyr
35	Asn	Thr 50	Asp	Thr	Asp	Asn	Cys 55	_	Glu	Val	Ile	Ile 60	Thr	Pro	Asn	Ser
40	Ala 65	Trp	Gly	Gly	Glu	Gly 70	Ser	Leu	Gly	Cys	Gly 75	Ile	Gly	Tyr	Gly	Туг 80
	Leu	His	Arg	Ile	Pro 85	Thr	Arg	Pro	Phe	Glu 90	Glu	Gly	Lys	Lys	Ile 95	Ser
45	Leu	Pro	Gly	Gln 100	Met	Ala	Gly	Thr	Pro 105	Ile	Thr	Pro	Leu	Lys 110	Asp	Gly
	Phe	Thr	Glu 115	Val	Gln	Leu	Ser	Ser 120	Val	Asn	Pro	Pro	Ser 125	Leu	Ser	Pro
50	Pro	Gly 130		Thr	Gly	Ile	Glu 135	Gln	Ser	Leu ·	Thr	Gly 140	Leu	Ser	Ile	Ser
55	Ser 145		•											٠		
	(2)	7370	OD\$45	m row	P~-	OT-0	TD	N7O	<b>51</b> F .							

(2) INFORMATION FOR SEQ ID NO: 515:

60 (i) SEQUENCE CHARACTERISTICS:

•				(	B) T	YPE:	ami	45 a no a	cid	aci	ds					
5			(xi)					lin PTIO		EQ I	D NO	: 51	5:		•	
3	Glu 1	Ser	Asn	Ser	Pro 5	Ala	Ala	Leu	Ala	Gly 10	Leu	Arg	Pro	His	Ser 15	Asp
10	Tyr	Ile	Ile	Gly 20	Ala	Asp	Thr	Val	Met 25	Asn	Glu	Ser	Glu	Asp 30	Leu	Phe
	Ser	Leu	Ile 35	Glu	Thr	His	Glu	Ala 40	Lys	Pro	Leu	Lys	Leu 45	Tyr	Val	Tyr
15	Asn	Thr 50	Asp	Thr	Asp	Asn	Cys 55	Arg	Glu	Val	Ile	Ile 60	Thr	Pro	Asn	Ser
20	Ala 65	Trp	Gly	Gly	Glu	Gly 70	Ser	Leu	Gly	Cys	Gly 75	Ile	Gly	Tyr	Gly	Tyr 80
	Leu	His	Arg	Ile	Pro 85	Thr	Arg	Pro	Phe	Glu 90	Glu	Gly	Lys	Lys	Ile 95	Ser
25	Leu	Pro	Gly	Gln 100	Met	Ala	Gly	Thr	Pro 105	Ile	Thr	Pro	Leu	Lys 110	Asp	Gly
	Phe	Thr	Glu 115	Val	Gln	Leu	Ser	Ser 120	Val	Asn	Pro	Pro	Ser 125	Leu	Ser	Pro
30	Pro	Gly 130	Thr	Thr	Gly	Ile	Glu 135	Gln	Ser	Leu	Thr	Gly 140	Leu	Ser	Ile	Ser
35	Ser 145													1		
	(2)	INF	ORMA:	rion	FOR	SEQ	ID	NO: 5	516:							
40		•	(i)	(	A) L B) T	ENGT YPE:	H: 1 ami	ERIS 51 a no a	mino cid		ds					
45			(xi)		-			lin PTIO		EQ I	OM C	: 51	6:			
	Arg 1	Ile	Pro	Thr	Arg 5	Pro	Phe	Glu	Glu	Gly 10	Lys	Lys	Ile	Ser	Leu 15	Pro
50	Gly	Gln	Met	Ala 20	Gly	Thr	Pro	Ile	Thr 25	Pro	Leu	Lys	Asp	Gly 30	Phe	Thr
	Glu	Val	Gln 35	Leu	Ser	Ser	Val	Asn 40	Pro	Pro	Ser	Leu	Ser 45	Pro	Pro	Gly
55	Thr	Thr 50	Gly	Ile	Glu	Gln	Ser 55	Leu	Thr	Gly	Leu	Ser 60	Ile	Ser	Ser	Thr
60	Pro 65	Pro	Ala	Val	Ser	Ser 70	Val	Leu	Ser	Thr	Gly 75	Val	Pro	Thr	Val	Pro 80

	Leu	Leu	Pro	Pro	Gln 85	Val	Asn	GIn	Ser	Leu 90	Thr	Ser	Val	Pro	95	Met
5	Asn	Pro	Ala	Thr 100	Thr	Leu	Pro	Gly	Leu 105	Met	Pro	Leu	Pro	Ala 110	Gly	Leu
	Pro	Asn	Leu 115	Pro	Asn	Leu	Asn	Leu 120	Asn	Leu	Pro	Ala	Pro 125	His	Ile	Met
10	Pro	Gly 130	Val	Gly	Leu	Pro	Glu 135	Leu	Val	Asn	Pro	Gly 140	Leu	Pro	Pro	Leu
15	Pro 145	Ser	Меt	Pro	Pro	Arg 150	Asn									
	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	VO: 5	517:							
20				(	A) L B) T D) T	CHAI ENGT YPE: OPOL	H: 1 ami OGY:	09 a no a lin	mino cid ear	aci			_			
25				_		E DE:				_					_	-1
	Pro 1	Gly	Leu	Pro	Pro 5	Leu	Pro	Ser	Met	Pro 10	Pro	Arg	Asn	Leu	Pro 15	GIY
30	Ile	Ala	Pro	Leu 20	Pro	Leu	Pro	Ser	Glu 25	Phe	Leu	Pro	Ser	Phe 30	Pro	Leu
	Val	Pro	Glu 35	Ser	Ser	Ser	Ala	Ala 40	Ser	Ser	Gly	Glu	Leu 45	Leu	Ser	Ser
35	Leu	Pro 50		Thr	Ser	Asn	Ala 55	Pro	Ser	Asp	Pro	Ala 60	Thr	Thr	Thr	Ala
40	Lys 65		Asp	Ala	Ala	Ser 70	Ser	Leu	Thr	Val	Asp 75	Val	Thr	Pro	Pro	Thr 80
	Ala	Lys	Ala	Pro	Thr 85	Thr	Val	Glu	Asp	Arg 90	Val	Gly	Asp	Ser	Thr 95	Pro
45	Val	Ser	Glu	Lys 100	Pro	Val	Ser	Ala	Ala 105		Asp	Ala	Asn			
50	(2)	INF				SEQ										
			(i)	(	(A) I (B) T	CHA ENGI TYPE:	H: 9	3 an ino a	nino cid		ls	•				
55			(xi)			OPOI E DE				EQ I	D NC	: 51	.8:			
	Ile 1	_	Lys	Val	Phe 5	_	His	Thr	Ala	Gly 10		Lys	Pro	Glu	Val 15	Ser
60	Cys	Phe	Glu	Asn	Ile	Arg	Ser	Cys	Ala	Arg	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa

				20	•				25					30		
5	Xaa	Xaa	Xaa 35	Xaa	Xaa	Xaa	Trp	Ile 40	Phe	Gly	Val	Leu	His 45	Val	Val	His
5	Ala	Ser 50	Val	Val	Thr	Ala	<b>T</b> yr 55	Leu	Phe	Thr	Val	Ser 60	Asn	Ala	Phe	Gln
10	Gly 65	Met	Phe	Ile	Phe	Leu 70	Phe	Leu	Cys	Val	Leu 75	Ser	Arg	Lys	Ile	Gln 80
	Glu	Glu	Tyr	Tyr	Arg 85	Leu	Phe	Lys	Asn	Val 90	Pro	Cys	Cys			
15						•				ě						
	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	: :OV	519:							
20				(	A) L B) T D) T	ENGT YPE: OPOL	H: 5 ami OGY:	5 am no a lin	ino cid ear	acid		: 51	9:			
25	Trp 1	Ile	Phe	Gly	Val 5	Leu	His	Val	Val	His 10	Ala	Ser	Val	Val	Thr 15	Ala
30	Tyr	Leu	Phe	Thr 20	Val	Ser	Asn	Ala	Phe 25	Gln	Gly	Met	Phe	Ile 30	Phe	Leu
50	Phe	Leu	Cys 35	Val	Leu	Ser	Arg	Lys 40	Ile	Gln	Glu	Glu	Тут 45	Tyr	Arg	Leu
35	Phe	Lys 50	Asn	Val	Pro	Cys	Cys 55									
40	(2)	INF			ENCE	CHA ENGI	RACT H: 5	ERIS 0 am	TICS		s					
45			(xi)		D) I	YPE: OPOL E DE	OGY:	lin	ear	EQ I	D NO	: 52	0:			
	Ala 1		Thr	Arg	Ile 5	Pro	Pro	Gly	Asp	Trp 10	Val	Ile	Asn	Val	Thr 15	Ala
50	Val	Ser	Phe	Ala 20	_	Lys	Thr	Thr	Ala 25	Arg	Phe	Phe	Xaa	His 30	Ser	Ser
55	Pro	Pro	Ser 35	Leu	Gly	Asp	Gln	Ala 40	_	Thr	Asp	Pro	Gly 45	His	Gln	Arg
	Arg	Asp 50														

PCT/US98/11422

	(2) INFORMATION FOR SEQ ID NO: 521:	
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 27 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 521:  Leu Gln Glu Val Asn Ile Thr Leu Pro Glu Asn Ser Val Trp Tyr Glu  1 5 10 15	
	Arg Tyr Lys Phe Asp Ile Pro Val Phe His Leu 20 25	
15		
	(2) INFORMATION FOR SEQ ID NO: 522:	
20	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 110 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 522:	
25	Met Gln Gly Ser Gly Ser Gln Phe Arg Ala Cys Leu Cys Leu Cys 1 5 10 15	
30	Phe Ser Cys Pro Cys Ser Pro Gly Gly Pro Arg Trp Asn Ser Arg Gln 20 25 30	
50	Gly Gly Arg Arg Phe Pro Lys Thr Cys Arg Ala Ile Ser Gln Asn Leu 35 40 45	
35	Val Phe Lys Tyr Lys Thr Phe Cys Pro Val Arg Tyr Met Gln Pro His 50 55 60	
	Arg Ser Ser Leu Cys Leu His Phe Thr Ser Tyr Val Phe Ile Leu Ser 65 70 75 80	
40	Thr Trp Gly Ser Leu Arg Thr Tyr Ser Thr Asp Leu Lys Lys Lys Lys 85 90 95	
45	Lys Asn Ser Arg Gly Gly Pro Val Pro Ile Arg Pro Lys Ser 100 105 110	
	(2) INFORMATION FOR SEQ ID NO: 523:	
50	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 99 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double	
55	(D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 523:	
	TAGCATGTAG CCAGTCGAAT AACNTATAAG GACAAAGTGG AGTCCACGCG TGCGGCCGTC	60
60	TAGACTAGTG GATCCCCCGG CTGCAGGATT CGGCACGAG	99

```
(2) INFORMATION FOR SEQ ID NO: 524:
5
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 51 amino acids
                    (B) TYPE: amino acid
10
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 524:
     Met Gln Gly Ser Gly Ser Gln Phe Arg Ala Cys Leu Leu Cys Leu Cys
15
      Phe Ser Cys Pro Cys Ser Pro Gly Gly Pro Arg Trp Asn Ser Arg Gln
      Gly Gly Arg Arg Phe Pro Lys Thr Cys Arg Ala Ile Ser Gln Asn Leu
20
                                  40
      Val Phe Lys
           50
25
     (2) INFORMATION FOR SEQ ID NO: 525:
              (i) SEQUENCE CHARACTERISTICS:
30
                     (A) LENGTH: 54 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 525:
      Pro Val Arg Tyr Met Gln Pro His Arg Ser Ser Leu Cys Leu His Phe
35
        1
      Thr Ser Tyr Val Phe Ile Leu Ser Thr Trp Gly Ser Leu Arg Thr Tyr
40
      Ser Thr Asp Leu Lys Lys Lys Lys Asn Ser Arg Gly Gly Pro Val
                                   40
       Pro Ile Arg Pro Lys Ser
45
           50
       (2) INFORMATION FOR SEQ ID NO: 526:
50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 38 amino acids
                      (B) TYPE: amino acid '
                     (D) TOPOLOGY: linear
 55
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 526:
       Gly Glu Glu Gln Arg Asp Cys Ser Leu Gly Trp Arg Gly Val Gly Met
                         5
                                            10
       Arg Ala Thr His Cys Gln Ala Ala Arg Met Phe Val Leu Phe Ser Leu
 60
```

25 30 20 Pro Lys Tyr Ala Gly Leu 35 5 (2) INFORMATION FOR SEQ ID NO: 527: (i) SEQUENCE CHARACTERISTICS: 10 (A) LENGTH: 161 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 527: 15 Met Pro Arg Lys Thr Ser Lys Cys Arg Gln Leu Leu Cys Ser Gly Ala Ser Arg Asn Ala Asp Thr Ala Ala Arg Gln Ser Thr Cys Ser Ser His 20 25 20 Arg Pro Pro Gly Lys Ile Pro Ser Leu Gly Pro Arg Arg Xaa Pro Gly Cys Xaa Ser Val Pro Ser Ser Arg Gly Glu Gln Ser Thr Gly Ser Pro 25 Ala Ala Pro Arg Cys Gly Arg Arg Asp Ala His Arg Gly Leu Pro Gly 70 30 Gly Ala Ala Met Thr Pro Gly Asp Thr Trp Ala Ser Phe Asn Pro Arg Ala Gly His Ser Lys Ser Gln Gly Glu Gly Gln Glu Ser Ser Gly Ala 35 Ser Arg Gln Asp Arg His Pro Val Ser His Trp Val Glu Arg Gln Arg 120 Glu Ala Trp Gly Ala Pro Arg Ser Ser Ser Ala Gly Gly Val Lys Val 40 135 Ala Ala Thr Thr Glu Arg Glu Pro Glu Phe Lys Ile Lys Thr Gly Lys 155 150 45 Ala 50 (2) INFORMATION FOR SEQ ID NO: 528: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 88 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 528:

Cys Ser Gly Ala Ser Arg Asn Ala Asp Thr Ala Ala Arg Gln Ser Thr

60

	Cys	Ser	Ser	His 20	Arg	Pro	Pro	Gly	Lys 25	Ile	Pro	Ser	Leu	Gly 30	Pro	Arg
5	Arg	Xaa	Pro 35	Gly	Cys	Xaa	Ser	Val 40	Pro	Ser	Ser	Arg	Gly 45	Glu	Gln	Ser -
10	Thr	Gly 50	Ser	Pro	Ala	Ala	Pro 55	Arg	Cys	Gly	Arg	Arg 60	Asp	Ala	His	Arg
	Gly 65	Leu	Pro	Gly	Gly	Ala 70	Ala	Met	Thr	Pro	Gly 75	Asp	Thr	Trp	Ala	Ser 80
15	Phe	Asn	Pro	Arg	Ala 85	Gly	His	Ser								
20	(2)	INF		TION SEOU												
25				~ (	A) L B) T D) T	ENGT YPE : OPOL	H: 5 ami OGY:	9 am no a lin	ino cid ear	acid		. 52	۵.			
23	Gln	Gly	•	SEQ Gly										Asp	Arg	His
30	1		50*	His	5	v-1	Gl.v	) ra	Cln	10	Glu	בומ	Tran	Glv	15	Pro
50	PIO	val	Ser	20	ırp	Val	Giu	n.y	25	ALG	GIU	AIG	TLD	30	ALG	110
35	Arg	Ser	Ser 35	Ser	Ala	Gly	Gly	Val 40		Val	Ala	Ala	Thr 45		Glu	Arg
	Glu	Pro 50		Phe	Lys	Ile	Lys 55		Gly	Lys	Ala					
40	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	530:							
45		•		1	(A) I (B) T (D) T	ENGT TYPE : TOPOI	TH: 2 ami	235 a ino a : lir	amino acid near	aci		): <b>5</b> 3	i <b>0</b> :			
50	Met		Pro	Arg	Tyr 5		Gly	Gly	Pro	Arg 10		Pro	Leu	Arg	Ile 15	Pro
	Asr	Glr	n Ala	Leu 20		Gly	Val	. Pro	Gly 25		Gln	Pro	Leu	Leu 30		Ser
55	Gly	/ Met	Asp 35		Thr	Arg	Glr	Glr 40		His	Pro	Asn	Met 45		Gly	Pro
60	Met	Glr 50		, Met	. Thr	Pro	Pro 55		, Gly	Met	. Val	Pro 60		Gly	Pro	Gln

	Asn 65	Tyr	Gly	Gly	Ala	Met 70	Arg	Pro	Pro	Leu	Asn 75	Ala	Leu	Gly	Gly	Pro 80
5	Gly	Met	Pro	Gly	Met 85	Asn	Met	Gly	Pro	Gly 90	Gly	Gly	Arg	Pro	Trp 95	Pro
	Asn	Pro	Thr	Asn 100	Ala	Asn	Ser	Ile	Pro 105	Tyr	Ser	Ser	Ala	Ser 110	Pro	Gly
10	Asn	Tyr	Val 115	Gly	Pro	Pro	Gly	Gly 120	Gly	Gly	Pro	Pro	Gly 125	Thr	Pro	Ile
15	Met	Pro 130	Ser	Pro	Ala	Asp	Ser 135	Thr	Asn	Ser	Gly	Asp 140	Asn	Met	Tyr	Thr
	Leu 145	Met	Asn	Ala	Val	Pro 150	Pro	Gly	Pro	Asn	Arg 155	Pro	Asn	Phe	Pro	Met 160
20	Gly	Pro	Gly	Ser	Asp 165		Pro	Met	Gly	Gly 170		Gly	Gly	Met	Glu 175	Ser
	His	His	Met	Asn 180		Ser	Leu	Gly	Ser 185	Gly	Asp	Met	Asp	Ser 190	Ile	Ser
25	Lys	Asn	Ser 195	Pro	Asn	. Asn	Met	Ser 200		Ser	Asn	Gln	Pro 205		Thr	Pro
30	Arg	Asp 210		Gly	Glu	ı Met	Gly 215		Asn	Phe	. Leu	Asn 220		Phe	Gln	Ser
50	Glu 225		Tyr	Ser	Pro	Ser 230		Thr	Met	Ser	Val 235					
35 ′	(2)	INI	FORM	AT I OI	1 FOI	R SEÇ	) ID	NO:	531 :							
			(i)	SEQ		E CHI										
40					(B)	TYPE TOPO	: am	ino	acid		103					
			(xi	) SE	QUEN	CE D				SEQ	ID N	o: 5	31:			
45		t Se	r Pr	o Ar		r. Pro 5	o Gly	y Gly	y Pro	Arg		Pro	Le	ı Ar	7 Ile 19	e Pro
	As	n Gl	n Al	a Le		y Gl	y Va	l Pro	o Gly 2		r Gli	n Pro	o Le	u Le		Ser
50	Gl	y Me		p Pr 5	o Th	r Ar	g Gl	n Gli		y Hi	s Pr	o Ası	n Me		y Gl	y Pro
e e	Ме		n Ar O	g Me	t Th	r Pr		o Ar 5	g Gl	y Me	t Va	l Pr		u Gl	y Pr	o Gln
55		n Ty 5	r Gl	y Gl	y Al		t Ar O	g Pr	o Pr	o Le	u As 7		a Le	u Gl	y Gl	y Pro 80
60	G1	у Ме	et Pr	. Gl		et As 35	n Me	t Gl	y Pr		y Gl O	y Gl	y Ar	g Pr	o Tr 9	p Pro 5

	Asn	Pro	Thr	Asn 100	Ala	Asn	Ser	Ile	Pro 105	Tyr	Ser	Ser	Ala	Ser 110	Pro	Gly
5	Asn	Tyr		٠												
10	(2)	INF	ORMA'	rion	FOR	SEQ	ID N	VO: 5	532:							
15				C	A) L B) T D) T	ENGT YPE: OPOL	H: 8 ami OGY:	1 am no a lin	ino cid ear	acid		: 53	2:			
20	Leu 1	Asn	Ala	Leu	Gly 5	Gly	Pro	Gly	Met	Pro 10	Gly	Met	Asn	Met	Gly 15	Pro
20	Gly	Gly	Gly	Arg 20	Pro	Trp	Pro	Asn	Pro 25	Thr	Asn	Ala	Asn	Ser 30	Ile	Pro
25	Tyr	Ser	Ser 35	Ala	Ser	Pro	Gly	Asn 40	Tyr	Val	Gly	Pro	Pro 45	Gly	Gly	Gly
	Gly	Pro 50		Gly	Thr	Pro	Ile 55	Met	Pro	Ser	Pro	Ala 60	Asp	Ser	Thr	Asn
30	Ser 65		y Asp	Asn	Met	<b>Tyr</b> 70	Thr	Leu	Met	Asn	Ala 75	Val	Pro	Pro	Gly	Pro 80
25	Asn															
35	(2)			m.r.o.v	<b>500</b>	. ~~~	<b>TD</b>	NO -	<b>533</b> .		•					
40 45	(2)	IN	(i)	1	ENCE (A) I (B) I	E CHA LENGI LYPE:	RACT	ERIS 70 an ino a : lir	TICS nino acid near	s: ació		): <b>5</b> 3	13: -	•		
43	Gly 1		o Met	Gly	Gly 5		Gly	Gly	Met	Glu 10		His	His	Met	Asn 15	
50	Ser	: Le	u Gly	y Ser 20		/ Asp	Met	. Asp	Ser 25		e Ser	Lys	. Asn	Ser 30		Asn
	Asr	n Me	t Sei 35	c Leu	Sei	: Asr	Glr	Pro 40		Thr	Pro	Arg	Asp 45		Gly	Glu
55	Met	: Gl; 5		y Asr	Phe	e Lev	ı Asr 55		Phe	e Glr	Sei	Glu 60		туг	Ser	Pro
60	Sei 65	_	t Thi	c Met	: Ser	val 70								•		

	(2) INFORMATION FOR SEQ ID NO: 534:
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 14 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 534:  Thr Cys Glu His Ser Ser Glu Ala Lys Ala Phe His Asp Tyr
	1 5 10
15	(2) INFORMATION FOR SEQ ID NO: 535:
20	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 59 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 535:
25	Gln Ala Phe Val Leu Leu Ser Asp Leu Leu Leu Ile Phe Ser Pro Gln 1 5 10 15
	Met Ile Val Gly Gly Arg Asp Phe Leu Arg Pro Leu Val Phe Pro 20 25 30
30	Glu Ala Thr Leu Gln Ser Glu Leu Ala Ser Phe Leu Met Asp His Val 35 40 45
35	Phe Ile Gln Pro Gly Asp Leu Gly Ser Gly Ala 50 55
	(2) INFORMATION FOR SEQ ID NO: 536:
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 43 amino acids  (B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 536:
45	Ala Cys Ser Tyr Leu Leu Cys Asn Pro Glu Phe Thr Phe Phe Ser Arg 1 5 10 15
50	Ala Asp Phe Ala Arg Ser Gln Leu Val Asp Leu Leu Thr Asp Arg Phe 20 25 30
	Gln Glu Leu Glu Glu Leu Leu Gln Val Gly 35 40
55	·
	(2) INFORMATION FOR SEQ ID NO: 537:
60	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 35 amino acids

```
(B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 537:
     Gln Lys Gln Leu Ser Ser Leu Arg Asp Arg Met Val Ala Phe Cys Glu
                . 5
     Leu Cys Gln Ser Cys Leu Ser Asp Val Asp Thr Glu Ile Gln Glu Gln
                                     25
10
      Val Ser Thr
15
      (2) INFORMATION FOR SEQ ID NO: 538:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 27 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 538:
      Gln Val Ile Leu Pro Ala Leu Thr Leu Val Tyr Phe Ser Ile Leu Trp
25
                                           10
      Thr Leu Thr His Ile Ser Lys Ser Asp Ala Ser
                                      25
                  20
30
      (2) INFORMATION FOR SEQ ID NO: 539:
             (i) SEQUENCE CHARACTERISTICS:
35
                    (A) LENGTH: 31 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 539:
40
      Ser Thr His Asp Leu Thr Arg Trp Glu Leu Tyr Glu Pro Cys Cys Gln
      Leu Leu Gln Lys Ala Val Asp Thr Gly Xaa Val Pro His Gln Val
                   20
45
      (2) INFORMATION FOR SEQ ID NO: 540:
50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 106 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 540:
55
      Leu Ala Val Ser Thr Ser Phe Ile Cys Cys Ala Asp Ile Ser Thr Ala
                                          10
      Leu Pro Leu Gly Ser Ser Arg Pro Ala Pro Ala Pro Arg His Arg Glu
60
```

	HIS	GIU	35	GIĀ	H1S	GIN	Ala	40	PIO	PIO	Arg	Leu	45	xaa	Thr	ser
5	Leu	Met 50	Pro	Leu	Ser	Thr	Pro 55	Ala	Ala	Ala	Gln	Leu 60	Leu	Trp	Thr	Gln
10	Leu 65	Thr	Pro	Met	Gly	Gly 70	Arg	Pro	Gly	Gly	Arg 75	His	Ser	Pro	Pro	Thr 80
	Leu	His	Thr	Gly	Pro 85	Arg	Ala	Leu	Pro	Pro 90	Gly	Pro	Pro	His	Pro 95	Ser
15	Leu	His	Val	Ala 100	Ala	Leu	Ser	Leu	Leu 105	Arg						
20	(2)	INFO	ORMAT			-				:						,
								07 a no a	mino cid	aci	ds					
25			(xi)					lin PTIO		EQ I	ои с	: 54:	1:			
	Glu 1	Gln	Val	Leu	Ala 5	Leu	Leu	Trp	Pro	Arg 10	Phe	Glu	Leu	Ile	Leu 15	Glu
. 30	Met	Asn	Val	Gln 20	Ser	Val	Arg	Ser	Thr 25	Asp	Pro	Gln	Arg	Leu 30	Gly	Gly
35	Leu	Asp	Thr 35	Arg	Pro	His	Tyr	Ile 40	Thr	Arg	Arg	Tyr	Ala 45	Glu	Phe	Ser
	Ser	Ala 50	Leu	Val	Ser	Ile	Asn 55	Gln	Thr	Ile	Pro	Asn 60	Glu	Arg	Thr	Met
40	Gln 65	Leu	Leu	Gly	Gln	Leu 70	Gln	Val	Glu	Val	Glu 75	Asn	Phe	Val	Leu	Arg 80
	Val	Ala	Ala	Glu	Phe 85	Ser	Ser	Arg	Lys	Glu 90	Gln	Leu	Val	Phe	Leu 95	Ile
45	Asn	Asn	Tyr	Asp 100	Met	Met	Leu	Gly	Val 105	Leu	Met	Glu	Arg	Ala 110	Ala	Asp
50	Asp	Ser	Lys 115	Glu	Val	Glu	Ser	Phe 120	Gln	Gln	Leu	Leu	Asn 125	Ala	Arg	Thr
, 50	Gln	Glu 130		Ile	Glu	Glu	Leu 135		Ser	Pro	Pro	Phe 140	Gly	Gly	Leu	Val
55	Ala 145	Phe	Val	Lys	Glu	Ala 150		Ala	Leu	Ile	Glu 155	Arg	Gly	Gln	Ala	Glu 160
	Arg	Leu	Arg	Gly	Glu 165		Ala	Arg	Val	Thr 170	Gln	Leu	Ile	Arg	Gly 175	Phe
60	Glv	Ser	Ser	Tro	Lvs	Ser	Ser	Val	Glu	Ser	Leu	Ser	Gln	Asp	Val	Met

,				180					185					190		
5	Arg	Ser	Phe 195	Thr	Asn	Phe	Arg	Asn 200	Gly	Thr	Ser	Ile	Ile 205	Gln	Gly	
	(2)	INF	ORMAT	rion	FOR	SEQ	ID N	10: 5	42:							
10			(i) :	(1	A) Li B) T	CHAR ENGTI YPE: OPOLO	d: 1	10 ar	mino cid		ds					
15			(xi)	SEQ						EQ II	ON 0	542	2:			
13	Ala 1		Leu	Lys	Tyr 5	Arg	Phe	Phe	Tyr	Gln 10	Phe	Leu	Leu	Gly	Asn 15	Glu
20	Arg	Ala	Thr	Ala 20	Lys	Glu	Ile	Arg	Asp 25	Glu	Tyr	Val	Glu	Thr 30	Leu	Ser
	Lys	Ile	Туr 35	Leu	Ser	Tyr	Тут	Arg 40	Ser	Tyr	Leu	Gly	Arg 45	Leu	Met	Lys
25 ،	Val	Gln 50	-	Glu	Glu	Val	Ala 55	Glu	Lys	Asp	Asp	Leu 60	Met	Gly	Val	Glu
30	Asp 65		Ala	Lys	Lys	Gly 70	Phe	Xaa	Ser	Lys	Pro 75	Ser	Leu	Arg	Ser	Arg 80
50	Asn	Thr	·Ile	Phe	Thr 85	Leu	Gly	Thr	Arg	Gly 90	Ser	Val	Ile	Ser	Pro 95	Thr
35	Glu	Leu	ı Glu	Ala 100	Pro	Ile	Leu	Val	Pro 105	His	Thr	Ala	Gln	Arg 110		•
40 .	(2)	INF		TION	ENCE	СНА	RACI	ERIS	TICS		ı.					
				(	(B) I	ENGI YPE:	ami	ino a	cid	acio	ıs					
45		,	(xi)	SEC		OPOL E DE				EQ I	D NO	: 54	3:			
	Glu 1		n Arg	ј Тут	Pro 5		Glu	Ala	Leu	Phe 10		Ser	Gln	His	Туг 15	Xaa
50	Leu	ı Lev	ı Asp	Asn 20		Cys	Arg	Glu	<b>Tyr</b> 25		Phe	Ile	Cys	Glu 30		Phe
55	Val	Va:	l Sex		Pro	Xaa	. Ala	His 40		Leu	Phe	His	Ala 45		Met	Gly
	Arg	7 Thi	_	ı Ser	Met	Thr	Leu 55		His	Leu	Asp	Ser 60		Leu	Ala	Asp
60	Cys 65		r Asp	) Ala	Ile	Ala		Phe	Leu	Cys	: Ile 75		Ile	val	Leu	Arg 80

	Phe Arg Asn Ile Ala Ala Lys Arg Asp Val Pro Ala Leu Asp Arg Tyr 85 90 95
5	Trp
10	(2) INFORMATION FOR SEQ ID NO: 544:
15	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 26 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 544:</li> </ul>
20	Gly Gly Leu Asp Thr Arg Pro His Tyr Ile Thr Arg Arg Tyr Ala Glu  1 5 10 15
	Phe Ser Ser Ala Leu Val Ser Ile Asn Gln 20 25
25	(2) INFORMATION FOR SEQ ID NO: 545:
30	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 20 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 545:</li> </ul>
35	Ser Arg Lys Glu Gln Leu Val Phe Leu Ile Asn Asn Tyr Asp Met Met  1 5 10 15
	Leu Gly Val Leu 20
40	(2) INFORMATION FOR SEQ ID NO: 546:
45	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 411 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 546:
50	Ala Leu Leu Lys Tyr Arg Phe Phe Tyr Gln Phe Leu Leu Gly Asn Glu 1 5 10 15
55	Arg Ala Thr Ala Lys Glu Ile Arg Asp Glu Tyr Val Glu Thr Leu Ser 20 25 30
<i>JJ</i>	Lys Ile Tyr Leu Ser Tyr Tyr Arg Ser Tyr Leu Gly Arg Leu Met Lys 35 40 45
60	Val Gln Tyr Glu Glu Val Ala Glu Lys Asp Asp Leu Met Gly Val Glu 50 55 60

	Asp 65	Thr	Ala	Lys	Lys	Gly 70	Phe	Xaa	Ser	Lys	<b>Pro</b> 75	Ser	Leu	-rg	Ser	Arg 80
5	Asn	Thr	Ile	Phe	Thr 85	Leu	Gly	Thr	Arg	Gl7 90	Ser	Val	Ile	Ser	95 95	Thr
	Glu	Leu	Glu	Ala 100	Pro	Ile	Leu	Val	Pro 105	His	Thr	Ala	Glm	Arg 110	Хаа	glu
10	Gln	Arg	Туг 115	Pro	Phe	Glu	Ala	Leu 120	Phe	Arg	Ser	Gln	His 125	Tyr	Хаа	Leu
15	Leu	Asp 130	Asn	Ser	Cys	Arg	Glu 135	Tyr	Leu	Phe	Ile	Cys 140	Glu	Phe	Phe	Val
	Val 145	Ser	Gly	Pro	Xaa	Ala 150	His	Asp	Leu	Phe	His 155	λίa	Val	Yet	GŢĀ	Arg 160
20	Thr	Leu	Ser	Met	Thr 165	Leu	Lys	His	Leu	Asp 170	Ser	TYT	Leu	Ala	Asp 175	Cys
25	Тут	Asp	Ala	Ile 180		Val	Phe	Leu	Cys 185	Ile	His	:ie	Va_	Leu 190	Arg	Phe
23	Arg	Asn	Ile 195		Ala	Lys	Arg	Asp 200	Val	Pro	Ala	Leu	Asp 205		Ty-z	طتن
30	Glu	Gln 210		Leu	Ala	Leu	Leu 215		Pro	Arg	Phe	Glu 220	Le:	lle	Leu	Glu
	Met 225		Val	Gln	Ser	Val 230		Ser	Thr	Asp	235		Arg	: Leu	Gly	31y 240
35	Leu	Asp	Thr	Arg	Pro 245		Tyr	Ile	Thr	Arg 250		Tyr	Ala	. Glu	255	: Ser
40	Ser	Ala	. Let	val 260		Ile	Asn	Gln	Thr 265		Pro	Asn	. Glu	<u>27</u> 9		Yet
10	Glr	Lev	1 Let 275		/ Gln	. Leu	Gln	Val 280		ı Val	Gl	i Ast	285		. Let	: Arg
45	Va]	Ala 290		a Glu	ı Phe	e Ser	Ser 295		Lys	Glu	ı Gl	1 Lev 300		: Phe	e Lev	: Ile
	Asr 305	_	а Туг	r Ası	) Met	310		ı Gly	Val	l Leu	1 Me: 31		ı Arş	ala Ala	a Als	320
50	Ası	Se:	r Ly:	s Glı	u Val 329	_	ı Ser	: Phe	e Gli	330		: Le	ı As:	n Ala	a Ar; 33:	Thr
55	Gli	n Gl	u Ph	e Il		ı Glu	ı Leı	ı Lev	34!		o Pr	es c	e Gl	y 31; 350	y Le	: Val
	Ala	a Ph	e Va 35		s Gl	u Ala	a Gl	u Ala 360		u Il	e Gl	u Ari	g Gl; 36	y Gli 5	n Al	a Glu
60 -	Ar	g Le		g Gl	y Gl	u Gl	u Ala 37		y Va	1 Th	r Gl	n Let 33	u Il 0	e Ar	g Gl;	y Phe

	Gly 385	Ser	Ser	Trp	Lys ·	Ser 390	Ser	Val	Glu	Ser	Leu 395	Ser	Gln	qzA	Val	Met 400
5	Arg	Ser	Phe	Thr	Asn 405	Phe 	Arg	Asn	Gly	Thr 410	Ser					
10	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	NO: 5	3 <b>4</b> 7:							
15				. (	A) L B) T D) T	ENGT YPE : OPOL	H: 3 ami OGY:	03 a no a line	mino cid ear	aci		: 54	7:			
20	Tyr 1	Glu	Gly	Lys	Glu 5	Phe	Asp	Tyr	Val	Phe 10	Ser	Ile	Asp	Val	Asn 15	Glu
	Gly	Gly	Pro	Ser 20	Tyr	Lys	Leu	Pro	Tyr 25	Asn	Thr	Ser	Asp	Asp 30	Pro	Trp
25	Leu	Thr	Ala 35	Tyr	Asn	Phe	Leu	Gln 40	Lys	Asn	Asp	Leu	Asn 45	Pro	Met	Phe
•	Leu	Asp 50	Gln	Val	Ala	Lys	Phe 55	Ile	Ile	Asp	Asn	Thr 60	Lys	Gly	Gln	Met
30	Leu 65	Gly	Leu	Gly	Asn	Pro 70	Ser	Phe	Ser	Asp	Pro 75	Phe	Thr	Gly	Gly	Gly 80
35	Arg	Tyr	Val	Pro	Gly 85	Ser	Ser	Gly	Ser	Ser 90	Asn	Thr	Leu	Pro	Thr 95	Ala
<i>33</i>	Asp	Pro	Phe	Thr 100	Gly	Ala	Gly	Arg	Tyr 105	Val	Pro	Gly	Ser	Ala 110	Ser	Met
40	Gly	Thr	Thr 115	Met	Ala	Gly	Val	Asp 120	Pro	Phe	Thr	Gly	Asn 125	Ser	Ala	Tyr
	Arg	Ser 130	Ala	Ala	Ser	Lys	Thr 135	Met	Asn	Ile	Tyr	Phe 140	Pro	Lys	Lys	Glu
45	Ala 145	Val	Thr	Phe	Asp	Gln 150	Ala	Asn	Pro	Thr	Gln 155	Ile	Leu	Gly	Lys	Leu 160
50	Lys	Glu	Leu	Asn	Gly 165	Thr	Ala	Pro	Glu	Glu 170	Lys	Lys	Leu	Thr	Glu 175	Asp
50	Asp	Leu	Ile	Leu 180	Leu	Glu	Lys	Ile	Leu 185		Leu	Ile	Cys	Asn 190	Ser	Ser
55	Ser	Glu	Lys 195		Thr	Val	Gln	Gln 200	Leu	Gln	Ile	Leu	Trp 205	Lys	Ala	Ile
	Asn	Cys 210		Glu	Asp	Ile	Val 215		Pro	Ala	Leu	Asp 220	Ile	Leu	Arg	Leu
60	Ser	Tle	Lva	Hie	Pro	Ser	Val	Δen	Glu	Asn	Phe	Cvs	Asn	Glu	ī.vs	Glu

	225	230	235	240
5	Gly Ala Gln Phe Sec. 24		Asn Leu Leu Asn Pro 250	Lys Gly 255
3	Lys Pro Ala Asn Gl 260	n Leu Leu Ala Leu 265	Arg Thr Phe Cys Asn 270	Cys Phe
10	Val Gly Gln Ala Gl 275	y Gln Lys Leu Met 280	Met Ser Gln Arg Glu 285	Ser Leu
	Met Ser His Ala Il 290	e Glu Leu Lys Ser 295	Gly Ser Asn Lys Asn 300	Ile
15				
	(2) INFORMATION FO	-		
20	(A) (B) (D)	TE CHARACTERISTICS LENGTH: 18 amino TYPE: amino acid TOPOLOGY: linear NCE DESCRIPTION: SI	acids	
25	His Ile Ala Leu Al	a Thr Leu Ala Leu 5	Asn Tyr Ser Val Cys 10	Phe His 15
	Lys Asp			
30				
	(2) INFORMATION FO	OR SEQ ID NO: 549:		
35	(A) (B)	CE CHARACTERISTICS LENGTH: 49 amino TYPE: amino acid TOPOLOGY: linear		
40		NCE DESCRIPTION: S	EQ ID NO: 549:	
40	His Asn Ile Glu G	ly Lys Ala Gln Cys 5	Leu Ser Leu Ile Ser 10	Thr Ile
45	Leu Glu Val Val G	ln Asp Leu Glu Ala 25	Thr Phe Arg Leu Leu 30	
	Leu Gly Thr Leu I 35	le Ser Asp Asp Ser 40	Asn Ala Val Gln Leu 45	Ala Lys
50	Ser			
55	(2) INFORMATION F	OR SEQ ID NO: 550:		
		ICE CHARACTERISTICS		•
		LENGTH: 30 amino TYPE: amino acid	acios	
60	(D)			

PCT/US98/11422

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 550:															
. 5	Leu 1	Gly	Val	Asp	Ser 5	Gln	Ile	Lys	Lys	Tyr 10	Ser	Ser	Val	Ser	Glu 15	Pro
J	Ala <sup>.</sup>	Lys	Val	Ser 20	Glu	Cys	Cys	Arg	Phe 25	Ile	Leu	Asn	Leu	Leu 30		
10	(2)	INFO	ORMAT	NOI	FOR	SEQ	ID N	ю: 5	51:							
15			(i) S (xi)	(1	A) Li B) T O) T	ENGT YPE : OPOL	H: 4 ami: OGY:	00 ar no ac line	mino cid ear	aci		: 551	l:			•
20	Tyr 1	Glu	Gly	Lys	Glu 5	Phe	Asp	Tyr	Val	Phe 10	Ser	Ile	Asp	Val	Asn 15	Glu
	Gly	Gly	Pro	Ser 20	Tyr	Lys	Leu	Pro	Tyr 25	Asn	Thr	Ser	Asp	Asp 30	Pro	Trp
25	Leu	Thr	Ala 35	Tyr	Asn	Phe	Leu	Gln 40	Lys	Asn	Asp	Leu	Asn 45	Pro	Met	Phe
30	Leu	Asp 50	Gln	Val	Ala	Lys	Phe 55	Ile	Ile	Asp	Asn	Thr 60	Lys	Gly	Gln	Met
50	Leu 65	Gly	Leu	Gly	Asn	Pro 70	Ser	Phe	Ser	Asp	Pro 75	Phe	Thr	Gly	Gly	Gly 80
35	Arg	Tyr	Val	Pro	Gly 85	Ser	Ser	Gly	Ser	Ser 90	Asn	Thr	Leu	Pro	Thr 95	Ala
,	Asp	Pro	Phe	Thr 100	Gly	Ala	Gly	Arg	Tyr 105	Val	Pro	Gly	Ser	Ala 110	Ser	Met
40	Gly	Thr	Thr 115	Met	Ala	Gly	Val	Asp 120	Pro	Phe	Thr	Gly	Asn 125		Ala	Tyr
45	Arg	Ser 130		Ala	Ser	Lys	Thr 135	Met	Asn	Ile	Tyr	Phe 140	Pro	Lys	Lys	Glu
	Ala 145		Thr	Phe	Asp	Gln 150		Asn	Pro	Thr	Gln 155	Ile	Leu	Gly	Lys	Leu 160
50	Lys	Glu	Leu	Asn	Gly 165	Thr	Ala	Pro	Glu	Glu 170		Lys	Leu	Thr	Glu 175	Asp
	Asp	Leu	lle	Leu 180		Glu	Lys	Ile	Leu 185		Leu	Ile	Cys	190		Ser
55	Ser	Glu	Lys 195		Thr	Val	Gln	Gln 200		Gln	Ile	Leu	Trp 205		Ala	Ile
60	Asn	210		Glu	Asp	Ile	Val 215		Pro	Ala	Leu	Asp 220		e Leu	Arg	Leu

	Ser 225	Ile	Lys	His	Pro	Ser 230	Val	Asn	Glu	Asn	Phe 235	Cys	Asn	Glu	Lys	Glu 240
5	Gly	Ala	Gln	.Phe	Ser 245	Ser	His	Leu	Ile	Asn 250	Leu	Leu	Asn	Pro	Lys 255	Gly
	Lys	Pro	Ala	Asn 260	Gln	Leu	Leu	Ala	Leu 265	Arg	Thr	Phe	Cys	Asn 270	Cys	Phe
10	Val	Gly	Gln 275	Ala	Gly	Gln	Lys	Leu 280	Met	Met	Ser	Gln	Arg 285	Glu	Ser	Leu
15	Met	Ser 290	His	Ala	Ile	Glu	Leu 295	Lys	Ser	Gly	Ser	Asn 300	Lys	Asn	Ile	His
-	Ile 305	Ala	Leu	Ala	Thr	Leu 310	Ala	Leu	Asn	Tyr	Ser 315	Val	Суз	Phe	His	Lys 320
20	Asp	His	Asn	Ile	Glu 325	Gly	Lys	Ala	Gln	Cys 330	Leu	Ser	Leu	Ile	Ser 335	Thr
	Ile	Leu	Glu	Val 340	Val	Gln	Asp	Leu	Glu 345	Ala	Thr	Phe	Arg	Leu 350	Leu	Val
25	Ala	Leu	Gly 355	Thr	Leu	Ile	Ser	Asp 360	Asp	Ser	Asn	Ala	Val 365	Gln	Leu	Ala
30	Lys	Ser 370	Leu	Gly	Val	Asp	Ser 375	Gln	Ile	Lys	Lys	Туг 380	Ser	Ser	Val	Ser
	Glu 385	Pro	Ala	Lys	Val	Ser 390	Glu	Cys	Cys	Arg	Phe 395	Ile	Leu	Asn	Leu	Leu 400
35																
40	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	552:							
.0			(i)		(A) I	ENG	H: 1	ERIS 139 a	mino		.ds					
45.			(xi)	SEC				lir PTIC		EQ I	D NC	: 55	2:			
	Tyr 1		Asn	Gln	Asp 5	Gly	Asp	Ile	Leu	Arg 10		Gln	Val	Leu	His 15	Glu
50	His	Ile	Gln	Arg 20		Ser	Lys	Val	Val 25		Ala	Asn	His	Arg 30		Leu
55	Gln	Ile	Pro 35		Val	Туг	Leu	Arg 40		Ala	. Pro	Trp	Pro 45		Ala	Gln
,,	Ser	Glu 50		e Arg	Thr	Ile	Ser 55		Туг	Lys	Thr	Pro 60		Asp	Lys	Val
60	Gln 65		: Ile	e Leu	Arg	Met 70		Ser	Thr	Ile	Met 75		Leu	. Lev	Ser	Leu 80

	Ala	Asn	Glu	Asp	Ser 85	Val	Pro	Gly		Asp 90	Asp	Phe	Val	Pro	Val 95	Leu
5	Val	Phe	Val	Leu 100	Ile	Lys	Ala	Asn	Pro 105	Pro	Cys	Leu	Leu	Ser 110	Thr	Val
10	Gln	Tyr	Ile 115	Ser	Ser	Phe	Тут	Ala 120	Ser	Cys	Leu	Ser	Gly 125	Glu	Glu	Ser
	Tyr	Trp 130	Trp	Met	Gln	Phe	Thr 135	Ala	Ala	Val	Glu					
15	(2)	·	ORMA'	TION	FOR	SEQ	ID I	NO: S	553 :						*	
20				(	A) L B) T D) T	ENGT YPE : OPOL	H: 1 ami OGY:	.44 a .no a lin	mino cid ear	aci		: 55	3:			
25 <sup>-</sup>	Tyr 1	Pro	Asn	Gln	Asp 5	Gly	Asp	Ile	Leu	Arg 10	Asp	Gln	Val	Leu	His 15	Glu
	His	Ile	Gln	Arg 20	Leu	Ser	Lys	Val	Val 25	Thr	Ala	Asn	His	Arg 30	Ala	Leu
30	Gln	Ile	Pro 35		Val	Tyr	Leu	Arg 40		Ala	Pro	Trp	Pro 45	Ser	Ala	Gln
35	Ser	G1u 50		Arg	Thr	Ile	Ser 55		Tyr	Lys	Thr	Pro 60		Asp	Lys	Val
33	Gln 65	-	Ile	e Leu	Arg	Met 70		Ser	Thr	Ile	Met 75		Leu	Leu	Ser	Leu 80
40	Ala	Asn	Glu	ı Asp	Ser 85		Pro	Gly	Ala	Asp 90		Phe	Val	Pro	Val 95	Leu
į	Val	. Phe	val	Leu 100		. Lys	Ala	Asn	Pro 105		Cys	Leu	Leu	Ser 110		Val
45	Gln	тут	: Ile 119		Ser	Phe	туг	120		Cys	Leu	Ser	Gly 125		Glu	Ser
50	Тух	Trg 130		Met	: Glr	Phe	135		Ala	Val	. Glu	Phe 140		: Lys	Thr	: Ile
55																

(2) INFORMATION FOR SEQ ID NO: 554:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 14 amino acids

60

(B) TYPE: amino acid

```
(D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 554:
      Tyr Pro Asn Gln Asp Gly Asp Ile Leu Arg Asp Gln Val Leu
 5
       (2) INFORMATION FOR SEQ ID NO: 555:
10
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 11 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
15
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 555:
      Glu Ala Pro Trp Pro Ser Ala Gln Ser Glu Ile
                        5
20
       (2) INFORMATION FOR SEQ ID NO: 556:
              (i) SEQUENCE CHARACTERISTICS:
25
                     (A) LENGTH: 21 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 556:
30
       Ser Gly Glu Glu Ser Tyr Trp Trp Met Gln Phe Thr Ala Ala Val Glu
                        5
       Phe Ile Lys Thr Ile
                   20
35
       (2) INFORMATION FOR SEQ ID NO: 557:
40
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 18 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 557:
45
      Ala Asp Asp Phe Val Pro Val Leu Val Phe Val Leu Ile Lys Ala Asn
                        5
                                            10
       Pro Pro
50
       (2) INFORMATION FOR SEQ ID NO: 558:
55
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 12 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
60
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 558:
```

WO 98/54963

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Tyr Lys Thr Pro Arg Asp Lys Val Gln Cys Ile Leu
                        5
 5
      (2) INFORMATION FOR SEQ ID NO: 559:
             (i) SEQUENCE CHARACTERISTICS:
10
                     (A) LENGTH: 15 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 559:
15
     Gly Ala Asp Asp Phe Val Pro Val Leu Val Phe Val Leu Ile Lys
                       5
20
      (2) INFORMATION FOR SEQ ID NO: 560:
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 12 amino acids
                     (B) TYPE: amino acid
25
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 560:
      Pro Val Leu Val Phe Val Leu Ile Lys Ala Asn Pro
                        5
30
      (2) INFORMATION FOR SEQ ID NO: 561:
35
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 17 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 561:
40
      Ser Ala Arg Ala Ser Thr Gln Pro Pro Ala Gly Gln His Pro Gly Pro
                       5
      Cys
45
      (2) INFORMATION FOR SEQ ID NO: 562:
50
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 33 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
55
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 562:
      Met Pro Gly Arg Trp Arg Trp Gln Arg Asp Met His Pro Ala Arg Lys
                        5
                                           10
                                                               15
60
     Leu Leu Ser Leu Leu Phe Leu Ile Leu Met Gly Thr Glu Leu Thr Gln
```

WO 98/54963 PCT/US98/11422

649

20 25 30 Asp 5 (2) INFORMATION FOR SEQ ID NO: 563: 10 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 19 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 563: \ 15 Ser Ala Ala Pro Asp Ser Leu Leu Arg Ser Ser Lys Gly Ser Thr Arg 10 5 Gly Ser Leu 20 (2) INFORMATION FOR SEQ ID NO: 564: 25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 564: Ala Ala Ile Val Ile Trp Arg Gly Lys Ser Glu Ser Arg Ile Ala Lys 5 1 35 Thr Pro Gly Ile 40 (2) INFORMATION FOR SEQ ID NO: 565: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 17 amino acids (B) TYPE: amino acid 45 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 565: Pro Leu Gly Ile Thr Leu Pro Leu Gly Ala Pro Glu Thr Gly Gly Gly 10 1 50 Asp 55 (2) INFORMATION FOR SEQ ID NO: 566: (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 20 amino acids

(B) TYPE: amino acid

(2) INFORMATION FOR SEQ ID NO: 570:

```
(D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 566:
     Cys Ala Ala Glu Thr Trp Lys Gly Ser Gln Arg Ala Gly Gln Leu Cys
5
     Ala Leu Leu Ala
10
      (2) INFORMATION FOR SEQ ID NO: 567:
             (i) SEQUENCE CHARACTERISTICS:
15
                    (A) LENGTH: 20 amino acids
                     (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 567:
      Phe Arg Gly Gly Thr Leu Val Leu Pro Pro Thr His Thr Pro Glu
20
      Trp Leu Ile Leu
25
      (2) INFORMATION FOR SEQ ID NO: 568:
30 ·
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 22 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 568:
35
      Met Arg Ser Ala Arg Pro Ser Leu Gly Cys Leu Pro Ser Trp Ala Phe
                                           10
      Ser Gln Ala Leu Asn Ile
40
                   20
      (2) INFORMATION FOR SEQ ID NO: 569:
45
            (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 22 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
50
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 569:
      Leu Leu Gly Leu Lys Gly Leu Ala Pro Ala Glu Ile Ser Ala Val Cys
                                            10
55
      Glu Lys Gly Asn Phe Asn
                   20
```

```
(i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 26 amino acids
                    (B) TYPE: amino acid
 5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 570:
     Val Ala His Gly Leu Ala Trp Ser Tyr Tyr Ile Gly Tyr Leu Arg Leu
10
      Ile Leu Pro Glu Leu Gln Ala Arg Ile Arg
                  20
15
      (2) INFORMATION FOR SEQ ID NO: 571:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 18 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 571:
      Thr Tyr Asn Gln His Tyr Asn Asn Leu Leu Arg Gly Ala Val Ser Gln
25
      Arg Cys
30
      (2) INFORMATION FOR SEQ ID NO: 572:
             (i) SEQUENCE CHARACTERISTICS:
35
                     (A) LENGTH: 43 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 572:
40
      Ile Leu Leu Pro Leu Asp Cys Gly Val Pro Asp Asn Leu Ser Met Ala
      Asp Pro Asn Ile Arg Phe Leu Asp Lys Leu Pro Gln Gln Thr Gly Asp
                   20
45
      Arg Ala Gly Ile Lys Asp Arg Val Tyr Ser Asn
50
       (2) INFORMATION FOR SEQ ID NO: 573:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 45 amino acids
55
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 573:
       Ser Ile Tyr Glu Leu Leu Glu Asn Gly Gln Arg Ala Gly Thr Cys Val
60
                                           10
                       5
        1
```

	Leu	Glu	Tyr	Ala 20	Thr	Pro	Leu	Gln	Thr 25	Leu	Phe	Ala	Met	Ser 30	Gln	Tyr
5	Ser	Gln	Ala 35	Gly	Phe	Ser	Gly	Glu 40	Asp	Arg	Leu	Glu	Gln 45			
10	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	NO: 5	574:							
15				(	A) Ļ B) T D) T	ENGT YPE : OPOL	H: 9 ami OGY:	2 am no a lin	ino cid ear	acid		: 57	4:			
20	Ala 1	Lys	Leu	Phe	Cys 5	Arg	Thr	Leu	Glu	Asp 10	Ile	Leu	Ala	Asp	Ala 15	Pro
20	Glu	Ser	Gln	Asn 20	Asn	Cys	Arg	Leu	Ile 25	Ala	Tyr	Gln	Glu	Pro 30	Ala	Asp
25	Asp	Ser	Ser 35	Phe	Ser	Leu	Ser	Gln 40	Glu	Val	Leu	Arg	His 45	Leu	Arg	Gln
	Glu	Glu 50	-	Glu	Glu	Val	Thr 55		Gly	Ser	Leu	Lys 60		Ser	Ala	Val
30	Pro 65		Thr	Ser	Thr	Met 70		Gln	Glu	Pro	Glu 75		Leu	Ile	Ser	Gly 80
35	Met	Glu	Lys	Pro	Leu 85		Leu	Arg	Thr	Asp 90		Ser				
40	(2)	INF		TION												
40			(1)		(A) 1 (B) 1	LENG: IYPE	TH: 4	43 ar ino a	mino acid		is					Ti.
45	Lov	. I o		) SE(	QUENC	CE DE	ESCRI	PTIC	ON: S					· Ala	Val	. Cys
	1	L			5	i			•	10					15	
50				20	· ·				25	5			ı TIÇ	30		Tyr
	Ile	e Gly	7 Ty: 3!	r Leu	ı Arç	j Let	ı Ile	Leu 40		o Glu	ı Lev	1				
55	(2)	) IN	FORM	ATIO	N FOI	R SE(	Q ID	NO:	576:	:						
60				SEQ	UENC:	E CH	ARAC		STIC	s:	ds					

```
(B) TYPE: amino acid
                    (D) TCPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 576:
 5
     Thr Met Lys Leu Lys Leu Arg Arg Asn Ile Val Lys Leu Ser Leu
           •
     Tyr Arg His Phe Thr Asn
10
      (2) DEFORMATION FOR SEQ ID NO: 577:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LEXGTH: 22 amino acids
                    (3) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 577:
20
     Thr Leu Ile Leu Ala Val Ala Ala Ser Ile Val Phe Ile Ile Trp Thr
                       5
       1
     Thr Met Lys Phe Arg Ile
25
                  20
      (2) DEFORMATION FOR SEQ ID NO: 578:
30
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 28 amino acids
                    'E) TYPE: amino acid
                    D, TOPOLOGY: linear
35
             (xii) SEQUENCE DESCRIPTION: SEQ ID NO: 578:
     Val Thr Cys Glm Ser Asp Trp Arg Glu Leu Trp Val Asp Asp Ala Ile
40
     Trp Arg Leu Leu Phe Ser Met Ile Leu Phe Val Ile
                  20
45
      (2) DIFGEMATION FOR SEQ ID NO: 579:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 27 amino acids
                    (3) TYPE: amino acid
50
                   (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 579:
     Met Val Leu Tro Arg Pro Ser Ala Asn Asn Gln Arg Phe Ala Phe Ser
                      · 5
55
      Pro Leu Ser Glu Glu Glu Glu Glu Asp Glu Gln
```

	(2) INFORMATION FOR SEQ ID NO: 580:
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 27 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 580:</li> </ul>
10	Met Val Leu Trp Arg Pro Ser Ala Asn Asn Gln Arg Phe Ala Phe Ser 1 5 10 15
	Pro Leu Ser Glu Glu Glu Glu Asp Glu Gln 20 25
15	•
	(2) INFORMATION FOR SEQ ID NO: 581:
20	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 35 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 581:</li> </ul>
25	Lys Glu Pro Met Leu Lys Glu Ser Phe Glu Gly Met Lys Met Arg Ser 1 5 10 15
30	Thr Lys Gln Glu Pro Asn Gly Asn Ser Lys Val Asn Lys Ala Gln Glu 20 25 30
	Asp Asp Leu 35
35	(2) INFORMATION FOR SEQ ID NO: 582:
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 37 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 582:
45	Lys Trp Val Glu Glu Asn Val Pro Ser Ser Val Thr Asp Val Ala Leu 1 5 10 15
	Pro Ala Leu Leu Asp Ser Asp Glu Glu Arg Met Ile Thr His Phe Glu 20 25 30
50	Arg Ser Lys Met Glu 35
55	(2) INFORMATION FOR SEQ ID NO: 583:
	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 20 amino acids</li><li>(B) TYPE: amino acid</li></ul>
60	(D) TOPOLOGY: linear

```
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 583:
      Asp Pro Arg Val Arg Leu Asn Ser Leu Thr Cys Lys His Ile Phe Ile
 5
      Ser Leu Thr Gln
10
      (2) INFORMATION FOR SEQ ID NO: 534:
            (i) SEQUENCE CERFACTERISTICS:
                    (A) LENGTH: 13 amino acids
15
                    (B) TYFE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 584:
      Tyr Glu Pro Met Asp Phe Maa Met Ala Lei Ile Tyr Asp
20
                       5
      (2) INFORMATION FOR SEQ ID NO: 535:
25
             (i) SEQUENCE CHAPACTERISTICS:
                    (A) LENGTH: 16 amino acids
                    (B) TFFE: amimo asid
                    (D) TOPOLOGY: linear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 515:
     Ile Arg His Glu Leu Thr Val Leu Arg Asp Thr Arg Fro Ala Dys Ala
35
40
      (2) INFORMATION FOR SEQ ID NO: 536:
          - (i) SEQUENCE THAFACTERISTICS:
                   (A) LENGTH: 10 amino acids
                    (B) TYPE: amimo asid
45
                    (D) TCPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 586:
      Met Asp Phe Xaa Met Ala Leu Ile Tyr Ast
          5
50
      (2) INFORMATION FOR SEQ ID NO: 587:
55
             (i) SEQUENCE CHAFACTERISTICS:
                    (A) LENGTH: 14 amino acids
                    (B) TIPE: amino asid
                  (D) TCPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 587:
60
```

•	Met 1	Gln	Glu	Met	Met 5	Arg	Asn	Gln	Asp	Arg 10	Ala	Leu	Ser	Asn	Leu 15	Glu
5	Ser	Ile	Pro	Gly 20	Gly	Tyr	Asn	Ala								
10	(2)	INF		(	ENCE A) L B) T	CHAI ENGT YPE:	RACT H: 2 ami	ERIS 5 am no a	PICS ino cid		s					
15			(xi)	SEQI		OPOL E DE:				EQ I	ON O	: 58	8:			
	Leu 1	Arg	Arg	Met	Tyr 5	Thr	Asp	Ile	Gln	Glu 10	Pro	Met	Leu	Ser	Ala 15	Ala
20	Gln	Glu	Gln	Phe 20	Gly	Gly	Asn	Pro	Phe 25							
25	(2)	INF	ORMA!	rion	FOR	SEQ	ID 1	vo: 5	89:							
30				(	A) L B) T D) T	ENGT YPE : OPOL	H: 3 ami OGY:	2 am no a lin	ino a cid ear	acid		: 58	9:			
35	Ala 1	Ser	Leu	Val	Ser 5	Asn	Thr	Ser	Ser	Gly 10	Glu	Gly	Ser	Gln	Pro 15	Ser
	Arg	Thr	Glu	Asn 20	Arg	Asp	Pro	Leu	Pro 25	Asn	Pro	Trp	Ala	Pro 30	Gln	Thr
40																
45	(2)	INF		(	ENCE A) L B) T		RACTI H: 7	ERIS 1 am no a	FICS: ino a		s					
50				SEQ	JENC	E DE	SCRI	PTIO	N: SI	-						
	Ser 1	Gln	Ser	Ser	Ser 5	Ala	Ser	Ser	Gly	Thr 10	Ala	Ser	Thr	Val	Gly 15	Gly
55	Thr	Thr	Gly	Ser 20	Thr	Ala	Ser	Gly	Thr 25	Ser	Gly	Gln	Ser	Thr 30	Thr	Ala
60	Pro	Asn	Leu 35	Val	Pro	Gly	Val	Gly 40	Ala	Ser	Met	Phe	Asn 45	Thr	Pro	Gly

WO 98/54963

	Met	Gln 50	Ser	Leu	Leu	Gln	Gln 55	Ile	Thr	Glu .	Asn	Pro 60	Gln	Leu	Met (	Gln
5	Asn 65	Met	Leu	Ser	Ala	Pro 70	Tyr									
10	(2)				ENCE	CHA	RACT	ERIS	591: TICS:		5					
15		•	(xi)	(	B) I D) I	YPE:	ami OGY:	no a	cid			: 59	1:			
	Met 1	Arg	Ser	Met	Met 5	Gln	Ser	Leu	Ser	Gln 10	Asn	Pro	Asp	Leu	Ala 15	Ala
20	Gln	Met	Met	Leu 20		Asn	Pro	Leu	Phe 25	Ala	Gly	Asn	Pro	Gln 30	Leu	Gln
25	Glu	Gln	Met 35		Gln	Gln	Leu	Pro 40	Thr	Phe	Leu	Gln	Gln 45			
	(2)	INF	ORMA	ATION	FOR	SEÇ	) ID	NO:	592:							
30			(i)		(A) : (B) :	LENG TYPE	TH: : am	73 a ino	STICS mino acid	: acid	ls					
35				) SE(	QUEN	CE D	ESCR		ON: S					•		<b>M</b>
	1					5		•		10	ı				15	
40				20	)				25	5				30	)	Glu
	Ala	Pro	Gl;		ıIl	e Pr	o Gl	y Ph		r Pro	Gly	/ Lev	1 Gly 45	/ Ala	a Leu	Gly
45	Ser	Th:		y Gly	y Se	r Se		y Th	r Ası	n Gly	/ Sei	ASI 6	n Ala	a Thi	Pro	Ser
50	Gli 6		n Th	r Se	r Pr	o Th 7		a Gl	y Th	r						
	(2	) İN	FORM	ATIO	N FC	R SE	Q II	NO:	593	:						
55			(i)	SEÇ	(A) (B)	LEN TYP	GTH: E; a	72 a mino	STIC amino ació inear	aci 1	.ds		-			
60			(x:	i) SE					ION:		ID N	iO: 5	93:			

	Thr (	3Lu	ޱo	Gly	His 5	Gln	Gln	Phe	Ile	Gln 10	Gln	Met	Leu	Gln	Ala 15	Leu
5	Ala :	31y	Val	Asn 20	Pro	Gln	Leu	Gln	Asn 25	Pro	Glu	Val	Arg	Phe 30	Gln	Gln
	ale :	Jeri	Glu 35	Gln	Leu	Ser	Ala	Met 40	Gly	Phe	Leu	Asn	Arg 45	Glu	Ala	Asn
10	Leu	مـــٰد 50	Ala	Leu	Ile	Ala	Thr 55	Gly	Gly	Asp	Ile	Asn 60	Ala	Ala	Ile	Glu
15	Arg 65	Leu	Leu	Gly	Ser	Gln 70		Ser								
20	(2)	그라		SEQU	FOR	CHA	RACI	TERIS	TICS	: acid	is					
25	Arg	Asn		SE	(B) (D) (D) (D) (D) (D) (D) (D) (D) (D) (D	ropoi E Di	LOGY ESCRI	: lir [PTIC	near N: S	. Met	: Arg	•		n Asp	Arg	, Ala
30	l Leu	Ser	: As:	ı Lei 20			: Ile	e Pro	Gly 25			c Asr	ı Ala	a Lev 30	15 1 Arg	g Arg
	Мес	בער	Thi 3		o Ile	e Glı	ı Glı	1 Pro 40		Le	ı Sei	r Ala	a Ala 4!	a 5		
35	(2)		FORM	ATIO:	N FO	R SE	δ IĎ	NO:	595	:						
40		•		*	(B)	LENC TYPE TOPO	TH: E: an OLOGY	13 a nino : li	mino acid near	aci l		ю: 5	95:			
45	Gly 1		n Pr	o Ph	e Al	a Se 5	r Le	u Va	l Se		n Th O	r Se	r Se	r		
50	·(2)	IN	FOR	ATIC	N FC	R SE	Q II	NO:	596	; :						
55					(B)	TYP:	GTH: E: a: OLOG	11 a mino Y: 1	amino acio inea	o ac d r		NO: !	596:			
60		u As 1	sn A	rg A	sp Pi	ro Le	eu P	co As	in Pi		rp A. 10	la				

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(2) INFORMATION FOR SEQ ID NO: 597:
5
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 17 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 597:
10
     Gly Lys Ile Leu Lys Asp Gln Asp Thr Leu Ser Gln His Gly Ile His
                       5
     Asp
15
      (2) INFORMATION FOR SEQ ID NO: 598:
20
           (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 14 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 598:
25
      Gly Leu Thr Val His Leu Val Ile Lys Thr Gln Asn Arg Pro
                       5
30
      (2) INFORMATION FOR SEQ ID NO: 599:
             (i) SEQUENCE CHARACTERISTICS:
35
                    (A) LENGTH: 18 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 599:
      Ser Glu Leu Gln Ser Gln Met Gln Arg Gln Leu Leu Ser Asn Pro Glu
40
                                          10
                      5
        1
      Met Met
45
       (2) INFORMATION FOR SEQ ID NO: 600:
50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 14 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 600:
 55
       Pro Glu Ile Ser His Met Leu Asn Asn Pro Asp Ile Met Arg
                                 10
                        5
```

```
(2) INFORMATION FOR SEQ ID NO: 501:
             (i) SEQUENCE CHAPACTERISTICS:
                    (A) LENGTH: 18 amino acids
                    (3) TYPE: amino acid
5
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 601:
     Arg Gln Leu Ile Met Ala Asn Pro Gln Met Gln Gln Leu Ile Gln Arg
10
             5
     Asn Pro
15
      (2) INFORMATION FOR SEQ ID NO: 502:
             (i) SEQUENCE CHAPACTERISTICS:
20
                    (A) LENGTH: 27 amino acids
                    (3) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 502:
      Asn Leu Cys. His Val Asp Cys Gln Asp Leu Leu Asn Pro Asn Leu Leu
25
                      5
       1
      Ala Gly Ile His Cys Ala Lys Arg Ile Val Ser
                   20
30
      (2) INFORMATION FOR SEQ ID NO: 503:
             (i) SEQUENCE CHAPACTERISTICS:
35
                    (A) LENGTH: 23 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 503:
40
      Leu Asp Gly Phe Glu Gly Tyr Ser Leu Ser Asp Trp Leu Cys Leu Ala
                               10
                       5
      Phe Val Glu Ser Lys Phe Asn
45
                   20
       (2) INFORMATION FOR SEQ ID NO: 604:
50
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 22 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
 55
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 604:
       Asn Glu Asn Ala Asp Gly Ser Phe Asp Tyr Gly Leu Phe Gln Ile Asn
                                         10
 60
      Ser His Tyr Trp Cys Asn
```

WO 98/54963

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(2) INFORMATION FOR SEQ ID NO: 605:
5
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 27 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
10
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 605:
      Asn Leu Cys His Val Asp Cys Gln Asp Leu Leu Asn Pro Asn Leu Leu
                              10
                       5
15
      Ala Gly Ile His Cys Ala Lys Arg Ile Val Ser
20
      (2) INFORMATION FOR SEQ ID NO: 606:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 13 amino acids
25
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 606:
      Ile Arg Glu Val Asn Glu Val Ile Gln Asn Pro Ala Thr
30
                       5
        1
      (2) INFORMATION FOR SEQ ID NO: 607:
35
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: - 30 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
40
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 607:
      Ile Thr Arg Ile Leu Leu Ser His Phe Asn Trp Asp Lys Glu Lys Leu
      Met Glu Arg Tyr Phe Asp Gly Asn Leu Glu Lys Leu Phe Ala
45
                   20
50
      (2) INFORMATION FOR SEQ ID NO: 608:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 23 amino acids
                     (B) TYPE: amino acid
 55
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 608:
      Asn Thr Arg Ser Ser Ala Gln Asp Met Pro Cys Gln Ile Cys Tyr Leu
                                           10
                        5
 60
```

```
Asn Tyr Pro Asn Ser Tyr Phe
20
```

5 (2) INFORMATION FOR SEQ ID NO: 609:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 60 amino acids
- (A) LENGIN: OU MILITO GER
- 10 (B) TYPE: amino acid
  - (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 609:
- Cys Asp Ile Leu Val Asp Asp Asn Thr Val Met Arg Leu Ile Thr Asp 15 1 5 10 15

Ser Lys Val Lys Leu Lys Tyr Gln His Leu Ile Thr Asn Ser Phe Val

20 Glu Cys Asn Arg Leu Leu Lys Trp Cys Pro Ala Pro Asp Cys His His 40 45

Val Val Lys Val Gln Tyr Pro Asp Ala Lys Pro Val 50 55 60

25

35

- (2) INFORMATION FOR SEQ ID NO: 610:
- 30 (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 52 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 610:

Cys Asp Ile Leu Val Asp Asp Asn Thr Val Met Arg Leu Ile Thr Asp

Ser Lys Val Lys Leu Lys Tyr Gln His Leu Ile Thr Asn Ser Phe Val
40 20 25 30

Glu Cys Asn Arg Leu Leu Lys Trp Cys Pro Ala Pro Asp Cys His His 35 40 45

- 45 Val Val Lys Val 50
- 50 (2) INFORMATION FOR SEQ ID NO: 611:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 60 amino acids
    - (B) TYPE: amino acid
- 55 (D) TOPOLOGY: linear
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 611:

Gly Cys Asn His Met Val Cys Arg Asn Gln Asn Cys Lys Ala Glu Phe 1 5 10 15

	Cys Trp Val Cys Leu Gly Pro Trp Glu Pro His Gly Ser Ala Trp Tyr 20 25 30
5	Asn Cys Asn Arg Tyr Asn Glu Asp Asp Ala Lys Ala Ala Arg Asp Ala 35 40 45
	Gln Glu Arg Ser Arg Ala Ala Leu Gln Arg Tyr Leu 50 55 60
10	
	(2) INFORMATION FOR SEQ ID NO: 612:
15	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 60 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 612:</li> </ul>
20	Phe Tyr Cys Asn Arg Tyr Met Asn His Met Gln Ser Leu Arg Phe Glu  1 5 10 15
25	His Lys Leu Tyr Ala Gln Val Lys Gln Lys Met Glu Glu Met Gln Gln 20 25 30
23	His Asn Met Ser Trp Ile Glu Val Gln Phe Leu Lys Lys Ala Val Asp 35 40 45
30	Val Leu Cys Gln Cys Arg Ala Thr Leu Met Tyr Thr 50 55 60
35	(2) INFORMATION FOR SEQ ID NO: 613:
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 60 amino acids  (B) TYPE: amino acid
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 613:
	Tyr Val Phe Ala Phe Tyr Leu Lys Lys Asn Asn Gln Ser Ile Ile Phe 1 5 10 15
45	Glu Asn Asn Gln Ala Asp Leu Glu Asn Ala Thr Glu Val Leu Ser Gly 20 25 30
50	Tyr Leu Glu Arg Asp Ile Ser Gln Asp Ser Leu Gln Asp Ile Lys Gln 35 40 45
	Lys Val Gln Asp Lys Tyr Arg Tyr Cys Glu Ser Arg 50 55 60
55	(2) INFORMATION FOR SEQ ID NO: 614:
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 37 amino acids
60	(B) TYPE: amino acid

	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 614:
5	Thr Gly Leu Glu Cys Gly His Lys Phe Cys Met Gln Cys Trp Ser Glu  1 5 10 15
	Tyr Leu Thr Thr Lys Ile Met Glu Glu Gly Met Gly Gln Thr Ile Ser 20 25 30
10	Cys Pro Ala His Gly 35
15	(2) INFORMATION FOR SEQ ID NO: 615:
20	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 21 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 615:</li> </ul>
	Met Trp Gly Tyr Leu Phe Val Asp Ala Ala Trp Asn Phe Leu Gly Cys
25	1 5 10 15
20	Leu Ile Cys Gly Trp 20
20	
30	(2) INFORMATION FOR SEQ ID NO: 616:
	(i) SEQUENCE CHARACTERISTICS:
25	(A) LENGTH: 46 amino acids (B) TYPE: amino acid
35	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 616:
40	Met His Phe Ile Ser Ser Gly Asn Val Ser Ala Ile Arg Ser Ser Ile 1 5 10 15
	Leu Leu Leu Arg Xaa Ser Leu Ser Tyr Leu Gly Asn Cys Leu Arg Val 20 25 30
45	Ser Ala Ile Phe Val Tyr Phe Leu Leu Phe Leu Leu Leu Ser 35 40 45
50	(2) INFORMATION FOR SEQ ID NO: 617:
	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 80 amino acids (B) TYPE: amino acid
55	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 617:
	Met Asp Gln Ala Leu Arg Gly Ser Pro Ser Glu Gly Phe Ser Thr Asp
60	1 5 10 15

	Pro	Ser	Pro	Pro 20	Gln	Val	Gly	Arg	Gln 25	Ile	Pro	Ser	Phe	Pro 30	Pro	Trp
5	Arg	Arg	Leu 35	Val	Leu	Pro	Lys	Ala 40	Ser	Gly	Cys	Phe	Leu 45	Glu	Arg	Glu
1	Trp	Trp 50	Leu	Cys	Val	Phe	Lys 55	Leu	Arg	Thr	Arg	Pro 60	Gly	Ala	Glu	Ala
10	His 65	Ala	Tyr	Asn	Ser	Ser 70	Ile	Leu	Gly	Gly	Arg 75	Gly	Lys	Gly	Ile	Thr 80
15																
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO:	618:							
20				(	A) L B) T D) T	ENGT YPE : OPOL	H: 1 ami OGY:	31 a no a lin	mino cid ear	aci						
25	•	•		SEQ										Pro	Glu	Gln
	1				5					10					15	
30				20					25	i				30		Val
	Glu	Phe	arg 35		Arg	Met	Glu	Lys 40		ı Val	Ser	Asp	Phe 45		Glr	Asp
35	Ser	Gl <sub>3</sub>		lle	: Lys	Lys	55 55		Glr	n Pro	Met	Asr 60		: Ile	: Glu	ı Arg
40	Ser 65		e Lev	His	: Asp	70		Glu	ı Val	l Ala	1 Gly 75		ı Thr	Ser	Phe	Ser 80
	Phe	e Gl	y Glu	ı Asp	Asp 85		Cy:	s Arg	TY1	r Va:		: Ile	e Phe	e Lys	9!	s Glu 5
45	Phe	e Al	a Pro	Ser 100		Glv	ı Gl	u Lei	ı Ası 10		с Тул	r Ar	g Arq	Gly 110	/ Gl: )	ı Glu
	Tr <u>r</u>	As	p Pro		ı Lys	s Ala	a Gl	u Glo 120		s Ar	g Ası	n Xa	a Ly: 12!	s Glu 5	ı Le	u Ala
50	Glı	n Ar 13	g Gli 0	n												

- 55 (2) INFORMATION FOR SEQ ID NO: 619:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 76 amino acids
    - (B) TYPE: amino acid
- 60 (D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 619:
e	Glu Glu Glu Ala Ala Gln Gln Gly Pro Val Val Val Ser Pro Ala Ser 1 5 10 15
5	Asp Tyr Lys Asp Lys Tyr Ser His Leu Ile Gly Lys Gly Ala Ala Lys 20 25 30
10	Asp Ala Ala His Met Leu Gln Ala Asn Lys Thr Tyr Gly Cys Xaa Pro 35 40 45
	Val Ala Asn Lys Arg Asp Thr Arg Ser Ile Glu Glu Ala Met Asn Glu 50 55 60
15	Ile Arg Ala Lys Lys Arg Leu Arg Gln Ser Gly Glu 65 70 75
20	(2) INFORMATION FOR SEQ ID NO: 620:
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 40 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 620:
30	Pro Pro Arg Arg Pro Ala Gln Leu Pro Leu Thr Pro Gly Ala Gly Gln 1 5 10 15
	Gly Ala Gly Arg Asp Lys Ala Ala Ala Ile Arg Ala His Pro Gly Ala 20 25 30
35	Pro Pro Leu Asn His Leu Leu Pro 35 40
40	(2) INFORMATION FOR SEQ ID NO: 621:  (i) SEQUENCE CHARACTERISTICS:
	<ul><li>(A) LENGTH: 28 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 621:  Ala Val Pro Gln Ala Gly Gly Lys Gln Val Phe Asp Leu Ser Pro Leu  1 5 10 15
50	1 5 10 13  Glu Leu Gly Tyr Val Arg Gly Met Cys Val Cys Val 20 25
55	(2) INFORMATION FOR SEQ ID NO: 622:
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 207 amino acids  (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi)	SECUENCE	DESCRIPTION:	SEQ	ID	NO:	622:
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Met Leu Pro Ala Leu Ala Ser Cys Cys His Phe Ser Pro Pro Glu Gln
1 5 10 15

5

Ala Ala Arg Leu Lys Lys Leu Gln Glu Gln Glu Lys Gln Gln Lys Val 20 25 30

Glu Phe Arg Lys Arg Met Glu Lys Glu Val Ser Asp Phe Ile Gln Asp 10 35 40 45

Ser Gly Gln Ile Lys Lys Phe Gln Pro Met Asn Lys Ile Glu Arg 50 55 60

Ser Ile Leu His Asp Val Val Glu Val Ala Gly Leu Thr Ser Phe Ser 65 70 75 80

Phe Gly Glu Asp Asp Asp Cys Arg Tyr Val Met Ile Phe Lys Lys Glu 85 90 95

20

Phe Ala Pro Ser Asp Glu Glu Leu Asp Ser Tyr Arg Arg Gly Glu Glu 100 105 110

Trp Asp Pro Gln Lys Ala Glu Glu Lys Arg Asn Xaa Lys Glu Leu Ala 25 115 120 125

Gln Arg Gln Glu Glu Glu Ala Ala Gln Gln Gly Pro Val Val Val Ser 130 135 140

Pro Ala Ser Asp Tyr Lys Asp Lys Tyr Ser His Leu Ile Gly Lys Gly 145 150 155 160

Ala Ala Lys Asp Ala Ala His Met Leu Gln Ala Asn Lys Thr Tyr Gly 165 170 175

Cys Xaa Pro Val Ala Asn Lys Arg Asp Thr Arg Ser Ile Glu Glu Ala 180 185 190

Met Asn Glu Ile Arg Ala Lys Lys Arg Leu Arg Gln Ser Gly Glu
40 195 200 205

(2) INFORMATION FOR SEQ ID NO: 623:

45

35

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 34 amino acids
  - (B) TYPE: amino acid
  - (D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 623:

Leu Leu Cys Pro Val Leu Asn Ser Gly Xaa Ser Trp Asn Phe Pro His 1 5 10 15

Pro Ser Gln Pro Glu Tyr Ser Phe His Gly Phe His Ser Thr Arg Leu 20 25 30

Trp Ile

	(2) INFORMATION FOR SEQ ID NO: 624:
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 28 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
.0	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 624:
	Pro Ser Thr Pro Trp Phe Leu Phe Leu Leu Gly Leu Thr Cys Pro Phe 1 5 10 15
15	Ser Thr Ser His Pro Arg Trp Asp Ser Ile Pro Pro 20 25
20	(2) INFORMATION FOR SEQ ID NO: 625:
-	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 227 amino acids
25	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 625:
23	Glu Leu Ser Ile Ser Ile Ser Asn Val Ala Leu Ala Asp Glu Gly Glu
	1 5 10 15
30	Tyr Thr Cys Ser Ile Phe Thr Met Pro Val Arg Thr Ala Lys Ser Leu 20 25 30
25.	Val Thr Val Leu Gly Ile Pro Gln Lys Pro Ile Ile Thr Gly Tyr Lys 35 40 45
35	Ser Ser Leu Arg Glu Lys Asp Thr Ala Thr Leu Asn Cys Gln Ser Ser 50 55 60
40	Gly Ser Lys Pro Ala Ala Arg Leu Thr Trp Arg Lys Gly Asp Gln Glu 65 70 75 80
	Leu His Gly Glu Pro Thr Arg Ile Gln Glu Asp Pro Asn Gly Lys Thr 85 90 95
45	Phe Thr Val Ser Ser Ser Val Thr Phe Gln Val Thr Arg Glu Asp Asp 100 105 110
	Gly Ala Ser Ile Val Cys Ser Val Asn His Glu Ser Leu Lys Gly Ala 115 120 125
50	Asp Arg Ser Thr Ser Gln Arg Ile Glu Val Leu Tyr Thr Pro Thr Ala 130 135 140
55	Met Ile Arg Pro Asp Pro Pro His Pro Arg Glu Gly Gln Lys Leu Leu 145 150 155 160
	Leu His Cys Glu Gly Arg Gly Asn Pro Val Pro Gln Gln Tyr Leu Try 165 170 175
40	Chu Luc Chu Chu Sor Wal Pro Pro Leu Lys Met Thr Gln Glu Ser Ald

WO 98/54963

55

60

Leu

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PCT/US98/11422

.

669

190 180 185 Leu Ile Phe Pro Phe Leu Asn Lys Ser Asp Ser Gly Thr Tyr Gly Cys 200 5 Thr Ala Thr Ser Asn Met Gly Ser Tyr Lys Ala Tyr Tyr Thr Leu Asn . 220 215 Val Asn Asp 10 225 (2) INFORMATION FOR SEQ ID NO: 626: 15 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 64 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 626: Glu Leu Ser Ile Ser Ile Ser Asn Val Ala Leu Ala Asp Glu Gly Glu 10 5 Tyr Thr Cys Ser Ile Phe Thr Met Pro Val Arg Thr Ala Lys Ser Leu 25 20 Val Thr Val Leu Gly Ile Pro Gln Lys Pro Ile Ile Thr Gly Tyr Lys 30 Ser Ser Leu Arg Glu Lys Asp Thr Ala Thr Leu Asn Cys Gln Ser Ser 35 (2) INFORMATION FOR SEQ ID NO: 627: 40 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 65 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 627: Cys Gln Ser Ser Gly Ser Lys Pro Ala Ala Arg Leu Thr Trp Arg Lys Gly Asp Gln Glu Leu His Gly Glu Pro Thr Arg Ile Gln Glu Asp Pro 50 20 Asn Gly Lys Thr Phe Thr Val Ser Ser Ser Val Thr Phe Gln Val Thr 35

Arg Glu Asp Asp Gly Ala Ser Ile Val Cys Ser Val Asn His Glu Ser

5	(2)	INFO	RMAT	NOI	FOR	SEQ	ID N	10: 6	28:							
5			(i) :		A) L	ENGT	H: 5	8 am	ino a	: acid	S					
10			(xi)	(	B) T D) T UENC	OPOL	OGY:	lin	ear	EQ II	OM C	: 621	3:			
	His 1	Glu	Ser	Leu	Lys 5	Gly	Ala	Asp	Arg	Ser 10	Thr	Ser	Gln	Arg	Ile 15	Glu
15	Val	Leu	Tyr	Thr 20	Pro	Thr	Ala	Met	Ile 25	Arg	Pro	Asp	Pro	Pro 30	His	Pro
20	Arg	Glu	Gly 35	Gln	Lys	Leu	Leu	Leu 40	His	Cys	Glu	Gly	Arg 45	Gly	Asn	Pro
	Val	Pro 50		Gln	Tyr	Leu	Trp 55	Glu	Lys	Glu						
25	(2)	INF	ORMA	TION	FOR	SEQ	ID I	NO:	629:							
30				(	(A) I (B) I (D) I	ENGT TYPE : TOPOI	TH: 5 ami LOGY:	ino a ino a : lir	nino ncid near	: acid		): 62	9:			
35	Trp		Lys	Glu	Gly 5		Val	Pro	Prò	Leu 10		Met	Thr	Gln	Glu 15	Ser
	Ala	Leu	ıle	Phe 20		Phe	. Lęu	. Asn	Lys 25		Asp	.Ser	Gly	Thr 30		Gly
40	Cys	Thr	: Ala		Ser	: Asr	Met	Gly 40		Tyr	Lys	· Ala	45		Thr	Leu
45	Asr	1 Va] 50		Asr												
	(2)	. IN	FORM	MOITA	I FOR	R SEC	) ID	NO:	630:	:						٠
50			(i)		(A) (B)	LENG	TH: : am	123 ino	amin acid	o ac	ids					
55			-		QUEN	CE D	ESCR	IPTI	ON:	SEQ :						
	:	1			!	5				10	)				19	
60	Gl	у Су:	s Al	a Len		y Ar	g Pr	o Gl	y Pho		Gl <sub>2</sub>	y Gl	y Pro	Thi 30		s Ser

	Gly	His	His 35	Lys	Ser	His	Pro	Gly 40	Pro	Ala	Gly	Gly	Asp 45	Tyr	Asn	Arg
5	Cys	Asp 50	Arg	Pro	Gly	Gln	Val 55	His	Leu	His	Asn	Pro 60	Arg	Gly	Thr	Gly
	Arg 65	Arg	Gly	Gln	Leu	His 70	Pro	Thr	Ala	Gly	Pro 75	Gly	Val	His	Arg	Arg 80
	Ala	Cys	Pro	Ser	Gln 85		Leu	Pro	His	Arg 90	Leu	Gly	Pro	Gly	Val 95	Pro
15	Cys	Pro	Ser	Pro 100	Ser	Leu	Thr	Pro	Val 105		Pro	Ser	Trp	Thr 110	Gln	Ser
	Trp	Cys	Gly 115	Leu	Pro	Gly	Тут	Thr 120		Ser	Ser					
20	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	631:			٠		٠,		
25					(A) 1 (B) 1 (D) 1	LENG IYPE IOPO	TH: : : a.m. LOGY	22 ar ino a : li	mino acid near	acio		o: 63	31:			
30	Val		s Glr	Leu	His		n Ala	a Val	Glr	1 Gly		s Ala	ı Lev	ı Gly	/ Arg	g Pro
35	Gly	y Phe	e Pro	Gl <u>y</u> 20		y Pro	<b>&gt;</b>									÷
	(2)	) IN		OITA												
40			(i)	SEQ	(A) (B)	LENG TYPE	TH: : an	TERI 42 a nino : li	mino acid	aci l	ds.		•			٠.
45			(xi	) SE							ID N	o: .6	32:			
45		o Th 1	r Hi	s Se		у Ні 5	s Hi	s Ly	s Se	r Hi 1		o Gl	y Pr	o Al	a Gl 1	y Gly 5
50	As	р Ту	r As	n Ar 2	_	s As	p Ar	g Pr		y Gl 5	n Va	l Hi	s Le	u Hi 3	s As O	n Pro
	Ar	g Gl		r Gl	y Ar	g Ar	g Gl		n Le 0	u Hi	s					,
55																
	(2	2) IN	FORM	TATIC	N FC	OR SE	Q II	NO:	633	:						
60			(i)	SEC				STERI			ids					

															•	
		(:	xi)		TOI	PE: 8 POLOC DESC	3Y: 1	linea	ar	Q ID	NO:	633 :				
5	Leu H	lis I	Proj	Thr A	Ala G	Sly F	ro G	ily V	al F	lis A 10	rg A	urg A	ala C	ys P	ro S 15	er
10	Gln (	3ln 1	Leu	Pro F 20	lis A	Arg I	Leu C	sly E	25	Sly N	/al E	Pro C	ys P	ro S 30	er F	ro
10	Ser I	Leu '	Thr 35	Pro V	Val I	Leu I	Pro S	Ser 7	rp :	Thr C	3ln S	Ser 1	rp C 45	ys G	ly I	eu
15	Pro (	51y 50	Tyr	Thr :	Ser S	Ser S	Ser 55									
20	(2)				NCE	CHAR INGTH	ACTE	RIST 6 am	ICS:	acid	ls					
25			(xi)		o) TO	PE: OPOLO E DES	ŒΥ:	line	ar	Q ID	NO:	634	:			
	Ser 1	Leu	Arg	Arg	Pro 5	Arg	Ser	Ala	Ala	Xaa 10	Gln	Thr	Leu '	Thr '	Thr 15	Phe
30	Leu	Ser	Ser	Val 20	Ser	Ser	Ala	Ser	Ser 25	Ser	Ala	Leu	Pro	Gly 30	Ser	Arg
35	Glu	Pro	Cys 35	Asp	Pro	Arg	Ala	Pro 40	Pro	Pro	Pro	Arg	Ser 45	Gly	Ser	Ala
, <b>, ,</b> ,	Ala	Ser 50	Cys	Cys	Ser	Cys	Cys 55	Cys	Ser	Cys	Pro	Arg 60	Arg	Arg	Ala	Pro
40	Leu 65	Arg	Ser	Pro	Arg	Gly 70	Ser	Lys	Arg	Arg	11e 75	Arg	Gln	Arg	Glu	Val 80
	Val	Asp	Lev	ı Tyr	Asn 85		Met	Cys	Leu	Gln 90	Gly	Pro	Ala	Gly	Val 95	Pro
45	Gly	Arg	Asg	Gly 100		Pro	Gly	Ala	Asn 105	Gly	Ile	Pro	Gly	Thr 110	Pro	Gly
50	Ile	Pro	Gl <sub>3</sub> 11	y Arg 5	Asp	Gly	Phe	Lys 120		Glu	Lys	Gly	Glu 125	Cys	Leu	Arg
30	Glu	Ser 130		e Glu	Glu	Ser	Trp 135		Pro	Asn	Tyr	Lys 140	Gln	Cys	Ser	Trp
55	Ser 145		r Le	u Asr	Туг	: Gly 150		Asp	Leu	Gly	Lys 155	Ile	Ala	Glu	Cys	Th:
	Phe	e Thi	r Ly	s Met	: Arg		Asn	Ser	Ala	Leu 170	Arg	Val	Leu	Phe	Ser	Gly

Ser Leu Arg Leu Lys Cys Arg Asn Ala Cys Cys Gln Arg Trp Tyr Phe

				180					185					190		
5	Thr	Phe	Asn 195	Gly	Ala	Glu	Cys	Ser 200	Gly	Pro	Leu	Pro	Ile 205	Glu	Ala	Ile
5	Ile	Туг 210	Leu	Asp	Gln	Gly	Ser 215	Pro	Glu	Met	Asn	Ser 220	Thr	Ile	Asn	Ile
10	His 225	Arg	Thr	Ser	Ser	Val 230	Glu	Gly	Leu	Cys	Glu 235	Gly	Ile	Gly	Ala	Gly 240
	Leu	Val	Asp	Val	Ala 245	Ile	Trp	Val	Gly	Thr 250	Cys	Ser	Asp	Tyr	Pro 255	Lys '
15	Gly	Asp	Ala	Ser 260	Thr	Gly	Trp	Asn	Ser 265	Val	Ser	Arg	Ile	Ile 270	Ile	Ġlu
20	Glu	Leu	Pro 275													
	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	635 :							
25			(i)		(A) 1 (B) 1	LENG: IYPE	rh: ( : am:	TERIS 51 ar	nino acid		ds					
30			(xi)					: lir [PTIC		EQ :	ID NO	o: 63	15:			
50	Ser 1		ı Arg	J Arg		Arg	g Ser	: Ala	Ala	10		Thr	Leu	Thr	Thr 15	Phe
35	Let	ı Sei	c Sei	: Val		c Sei	Ala	a Sei	: Ser 25		c Ala	a Lev	Pro	Gl <sub>y</sub> 30	/ Sei	Arg
	Gl	ı Pro	o Cy:		) Pro	o Arg	Ala	a Pro		Pro	o Pro	o Arg	Sei 45	c Gly	/ Sei	c Ala
40	Ala	a Se:		s Cy:	s Se	r Cyr	s Cy: 5	s Cy: 5	s Se	r Cys	s Pro	Arg 60		3		
45	(2	) IN						NO:								
			(i)	SEQ	(A)	LENC	TH:	TERI 52 a	mino	aci	lds			•		
50			(xi	.) SE	(D)	TOPO	LOG	nino (: li RIPTI	near	:	ID N	10: 6	36:			
55		1				5				1	.0				1	g Gln 5
32	Ar	g Gl	u Va		ll As	sp L∈	eu Ty	n As		y M∈ !5	et Cy	rs L∈	u Gl	n Gl	y Pr 0	o Ala
-60	G]	ly Va		ro GI	.y Aı	g As	sp G		er Pr	:o G1	ly Al	a As	n Gl	ly II 15	.e Pr	o Gly

```
Thr Pro Gly Ile
          50
5
      (2) INFORMATION FOR SEQ ID NO: 637:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 52 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 637:
      Thr Pro Gly Ile Pro Gly Arg Asp Gly Phe Lys Gly Glu Lys Gly Glu
15
      Cys Leu Arg Glu Ser Phe Glu Glu Ser Trp Thr Pro Asn Tyr Lys Gln
20 -
      Cys Ser Trp Ser Ser Leu Asn Tyr Gly Ile Asp Leu Gly Lys Ile Ala
                                   40
      Glu Cys Thr Phe
25
           50
      (2) INFORMATION FOR SEQ ID NO: 638:
30
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 66 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 638:
35
      Phe Thr Lys Met Arg Ser Asn Ser Ala Leu Arg Val Leu Phe Ser Gly
       Ser Leu Arg Leu Lys Cys Arg Asn Ala Cys Cys Gln Arg Trp Tyr Phe
40
                                        25
       Thr Phe Asn Gly Ala Glu Cys Ser Gly Pro Leu Pro Ile Glu Ala Ile
                                    40
45
       Ile Tyr Leu Asp Gln Gly Ser Pro Glu Met Asn Ser Thr Ile Asn Ile
       His Arq
 50
        65
       (2) INFORMATION FOR SEQ ID NO: 639:
 55
               (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 51 amino acids
```

(B) TYPE: amino acid(D) TOPOLOGY: linear

60

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 639:

	Arg Thr Ser Ser Val Glu Gly Leu Cys	Glu Gly Ile Gly Ala Gly Leu 10 15
5	Val Asp Val Ala Ile Trp Val Gly Thr 20 25	
10	Asp Ala Ser Thr Gly Trp Asn Ser Val	Ser Arg Ile Ile Ile Glu Glu 45
	Leu Pro Lys 50	
15	(2) INFORMATION FOR SEQ ID NO: 640	· .
20	(i) SEQUENCE CHARACTERISTIC  (A) LENGTH: 26 amino  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION:	acids !
25	Thr Lys Lys Glu Asn Cys Arg Pro Al.	a Ser Leu Met Asn Ile Asp Thr 10 15
	Lys Ile Leu Asn Lys Ile Leu Met As 20 2	
30		
35	(2) INFORMATION FOR SEQ ID NO: 641  (i) SEQUENCE CHARACTERISTIC  (A) LENGTH: 214 amin  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION:	CS: no acids d r
40	Met Cys Asn Leu Pro Ile Lys Val Va	al Cys Arg Ala Asn Ala Glu Tyr 10 15
٠.	Met Ser Pro Ser Gly Lys Val Pro Xa 20	aa Xaa His Val Gly Asn Gln Val 25 30
45	Val Ser Glu Leu Gly Pro Ile Val G	ln Phe Val Lys Ala Lys Gly His 45
50	Ser Leu Ser Asp Gly Leu Glu Glu Vo	al Gln Lys Ala Glu Met Lys Ala 60
-	Tyr Met Glu Leu Val Asn Asn Met L 65 70	eu Leu Thr Ala Glu Leu Tyr Leu 75 80
55	Gln Trp Cys Asp Glu Ala Thr Val G 85	ly Xaa Ile Thr His Xaa Arg Tyr 90 95
	Gly Ser Pro Tyr Pro Trp Pro Leu X	aa His Ile Leu Ala Tyr Gln Lys 05 110

	Gln T		Glu 115	Val	Lys	Arg	Lys	Хаа 120	Lys	Ala	Ile	Gly	Trp 125	Gly	Lys	Lys
5	Thr L	eu 30	Asp	Gln	Val	Leu	Glu 135	Asp	Val	Asp	Gln	Cys 140	Cys	Gln	Ala	Leu
	Ser G 145	ln	Arg	Leu	Gly	Thr 150	Gln	Pro	Tyr	Phe	Phe 155	Asn	Lys	Gln	Pro	Thr 160
10	Glu L	eu	Asp	Ala	Leu 165	Val	Phe	Gly	His	Leu 170	Tyr	Thr	Ile	Leu	Thr 175	Thr
15	Gln L	eu	Thr	Asn 180	Asp	Glu	Leu	Ser	Glu 185	Lys	Val	Lys	Asn	Tyr 190	Ser	Asn
13	Leu I	eu	Ala 195	Phe	Cys	Arg	Arg	Ile 200		Gln	His	Tyr	Phe 205	Glu	Asp	Arg
20	Gly I	Lys 210	Gly	Arg	Leu	Ser				÷						
25	(2)								642 :							
			(1)	. (	(A) I	ENG	TH: 4		nino		ls					
30			(xi)		(D) 1	ropoi	LOGY	: lir		EQ I	D NO	): 64	12:			
	Met (	Cys	Asr	. Leu	Pro		Lys	. Val	. Val	Cys		Ala	. Asr	Ala	Glu 15	Tyr
35	Met	Ser	Pro	Ser 20		, Lys	val	l Pro	Хаа 25		. His	Va]	Gly	Asr 30		ı Val
40	Val	Ser	Glu 35		ı Gİy	, Pro	o Ile	e Val 40		n Phe	e Val	Lys	3	•		
40			35	5				40	)		e Val	L Lys	3			
40 45			35 ORM	5 ATION	N FOI	R SE( E CH	Q ID	40 NO: TERI	643: STIC:	: S:		L Lys	5	,		
			39 ORM (i)	ATION SEQ	N FOR UENC (A) (B) (D)	R SEC E CH LENG TYPE TOPO	Q ID  ARAC  TH:  : an	NO: TERI 44 a ino : li	643: STIC: mino acid near	s: aci	ds					
	(2)	INF	ORM (i)	ATION SEQ	N FOR UENC: (A) (B) (D) QUEN a Ly	R SEC E CH LENG TYPE TOPC CE D	Q ID ARAC TH: : am ILOGY ESCR	NO: TERI 44 a ino : li IPTI	643: STIC: mino acid near ON:	s: aci SEQ u Se	ds ID N	o: 6	43:	u Gl		u Val
45	(2) Phe 1	INF	ORMA (i) (xi	SEQ ) SE s Al	N FOR UENC (A) (B) (D) QUEN a Ly	R SEG E CH LENG TYPE TOPO CE D S G1;	Q ID  ARAC  TH:  : an  LOGY ESCR	NO: TERI 44 a ino 7: li IPTI s Se	643: STIC: mino acid near ON: r Le	S: aci SEQ u Se 1	ds ID N r As 0	O: 6	43: y Le		u Gl 1	u Val 5 t Leu

	(2) INFORMATION FOR SEQ ID NO: 644:
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 51 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 644:</li> </ul>
10	Leu Gln Trp Cys Asp Glu Ala Thr Val Gly Xaa Ile Thr His Xaa Arg 1 5 10 15
1.5	Tyr Gly Ser Pro Tyr Pro Trp Pro Leu Xaa His Ile Leu Ala Tyr Gln 20 25 30
15	Lys Gln Trp Glu Val Lys Arg Lys Xaa Lys Ala Ile Gly Trp Gly Lys 35 40 45
20	Lys Thr Leu 50
25	(2) INFORMATION FOR SEQ ID NO: 645:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 43 amino acids  (B) TYPE: amino acid
30	(D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 645:
	Asp Gln Val Leu Glu Asp Val Asp Gln Cys Cys Gln Ala Leu Ser Gln 1 5 10 15
35	Arg Leu Gly Thr Gln Pro Tyr Phe Phe Asn Lys Gln Pro Thr Glu Leu 20 25 30
40	Asp Ala Leu Val Phe Gly His Leu Tyr Thr Ile 35 40
45	(2) INFORMATION FOR SEQ ID NO: 646:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 41 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 646:  Leu Thr Thr Gln Leu Thr Asn Asp Glu Leu Ser Glu Lys Val Lys Ass  1 5 10 15
55	Tyr Ser Asn Leu Leu Ala Phe Cys Arg Arg Ile Glu Gln His Tyr Ph 20 25 30
	Glu Asp Arg Gly Lys Gly Arg Leu Ser 35 40

	(2) INFORMATION FOR SEQ ID NO: 647:
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 70 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 647:</li> </ul>
10	Met Xaa Xaa Xaa Asn Ser His Ile Thr Ile Fhe Thr Leu Asn Val Asn 1 5 10 15
	Gly Leu Asn Ala Pro Asn Glu Arg His Arg Leu Ala Asn Trp Ile Gln 20 25 30
15	Ser Gln Asp Gln Val Cys Cys Ile Gln Glu Thr His Leu Thr Gly Arg
20	Asp Thr His Arg Leu Lys Ile Lys Gly Trp Arg Lys Ile Tyr Gln Ala 50 55 60
	Asn Gly Lys Gln Lys Lys 65 70
25	
	(2) INFORMATION FOR SEQ ID NO: 648:
30	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 28 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 648:</li> </ul>
35	Phe Thr Leu Asn Val Asn Gly Leu Asn Ala Pro Asn Glu Arg His Arg 1 5 10 15
40	Leu Ala Asn Trp Ile Gln Ser Gln Asp Gln Val Cys 20 25
.0	
	(2) INFORMATION FOR SEQ ID NO: 649:
45	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 17 amino acids</li><li>(B) TYPE: amino acid</li></ul>
50	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 649:
	Thr His Leu Thr Gly Arg Asp Thr His Arg Leu Lys Ile Lys Gly Tr 1 5 10 15
55	Arg
	•
60	(2) INFORMATION FOR SEQ ID NO: 650:

	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 14 amino acids
	(B) TYPE: amino acid
_	(D) TOPOLOGY: linear
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 650:
•	Clumber Aver Inc. The Mer Cla Ale Aca Cly Ive Cla Ive Ive
	Gly Trp Arg Lys Ile Tyr Gln Ala Asn Gly Lys Gln Lys Lys 1 5 10
	1 3
10	
10	
	(2) INFORMATION FOR SEQ ID NO: 651:
	_
	(i) SEQUENCE CHARACTERISTICS:
15	(A) LENGTH: 54 amino acids
	(B) TYPE: amino acid
	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 651:
20	The stire to the Government of the Discount of
20	Ile Tyr His Leu His Ser Trp Ile Phe Phe His Phe Lys Arg Ala Phe
	1 5 10 15
	Cys Met Cys Phe Ile Thr Met Lys Val Ile His Ala His Cys Ser Lys
	20 25 30
25	20
	Leu Arg Lys Cys Xaa Asn Ala Gln Ile Ser Val Phe Cys Thr Thr Leu
	35 40 45
	Thr Ala Ser Tyr Pro Thr
30	50
	(2) INFORMATION FOR SEQ ID NO: 652:
35	(2) INFORMATION FOR SEQ 10 NO. 032.
JJ	(i) SEQUENCE CHARACTERÍSTICS:
	(A) LENGTH: 23 amino acids
	(B) TYPE: amino acid
	(D) TOPOLOGY: linear
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 652:
	Ile Tyr His Leu His Ser Trp Ile Phe Phe His Phe Lys Arg Ala Phe
	1 5 10 15
45	A A A A A A A A A A A A A A A A A A A
43	Cys Met Cys Phe Ile Thr Met
	20
50	(2) INFORMATION FOR SEQ ID NO: 653:
	- · ·
	(i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 31 amino acids
<b></b> -	(B) TYPE: amino acid
55	(D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 653:
	The stall the stall the Good Control Law Same Law Chan Very Same Stall
	Lys Val Ile His Ala His Cys Ser Lys Leu Arg Lys Cys Xaa Asn Ala 1 5 10 15
60	1 5 10 15
-	

PCT/US98/11422

680

Gln Ile Ser Val Phe Cys Thr Thr Leu Thr Ala Ser Tyr Pro Thr 25 20 5 (2) INFORMATION FOR SEQ ID NO: 654: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 58 amino acids 10 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 654: Trp Asn Leu Leu Trp Tyr Phe Gln Arg Leu Arg Leu Pro Ser Ile Leu 15 Pro Gly Leu Val Leu Ala Ser Cys Asp Gly Pro Ser Xaa Ser Gln Ala 20 Pro Ser Pro Trp Leu Thr Pro Asp Pro Ala Ser Val Gln Val Arg Leu 40 Leu Trp Asp Val Leu Thr Pro Asp Pro Asn 50 25 (2) INFORMATION FOR SEQ ID NO: 655: 30 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 54 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 655: 35 Gln Arg Gly Ile Tyr Arg Glu Ile Leu Phe Leu Thr Met Ala Ala Leu Gly Lys Asp His Val Asp Ile Val Ala Phe Asp Lys Lys Tyr Lys Ser 40 Ala Phe Asn Lys Leu Ala Ser Ser Met Gly Lys Glu Glu Leu Arg His 40 45 Arg Arg Ala Gln Met Pro 50 50 (2) INFORMATION FOR SEQ ID NO: 656: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 amino acids (B) TYPE: amino acid 55 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 656: Trp Asn Leu Leu Trp Tyr Phe Gln Arg Leu Arg Leu Pro Ser Ile Leu 60

Pro Gly Leu Val Leu Ala Ser 20

5	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: (	557 :							
10	,_,		(i) (xi)	SEQU ( ) (	ENCE A) L B) T D) T	CHA ENGT YPE: OPOL	RACT H: 1 ami OGY:	ERIS 91 a no a lin	TICS mino cid ear	aci	•	: 65	7:			
15	Glu 1	Asp	Asp	Gly	Phe 5	Asn	Arg	Ser	Ile	His 10	Glu	Val	Ile	Leu	Lys 15	Asn
	Ile	Thr	Trp	Туг 20	Ser	Glu	Arg	Val	Leu 25	Thr	Glu	Ile	Ser	Leu 30	Gly	Ser
20	Leu	Leu	Ile 35	Leu	Val	Val	Ile	Arg 40	Thr	Ile	Gln	Тут	Asn 45	Met	Thr	Arg
25	Thr	Arg 50	Asp	Lys	Tyr	Leu	His 55	Thr	Asn	Cys	Leu	Ala 60	Ala	Leu	Ala	Asn
	Met 65	Ser	Ala	Gln	Phe	Arg 70	Ser	Leu	His	Gln	Туг 75	Ala	Ala	Gln	Arg	Ile 80
80	Ile	Ser	Leu	Phe	Ser 85	Leu	Leu	Ser	Lys	Lys 90	His	Asn	Lys	Val	Leu 95	Glu
	Gln	Ala	Thr	Gln 100	Ser	Leu	Arg	Gly	Ser 105	Leu	Ser	Ser		Asp 110	Vaļ	Pro
35	Leu	Pro	Asp 115	Tyr	Ala	Gln	Asp	Leu 120	Asn	Val	Ile	Glu	Glu 125	Val	Ile	Arg
	Met	Met 130	Leu	Glu	Ile	Ile	Asn 135	Ser	Cys	Leu	Thr	Asn 140	Ser	Leu	His	His
	Asn 145	Pro	Asn	Leu	Val	Туг 150	Ala	Leu	Leu	Tyr	Lys 155	Arg	Asp	Leu	Phe	Glu 160
15	Gln	Phe	Arg		His 165	Pro	Ser	Phe	Gln	Asp 170	Ile	Met	Gln	Asn	Ile 175	Asp
	Leu	Val	Ile	Ser 180	Phe	Phe	Ser	Ser	Arg 185	Leu	Leu	Gln	Ala	Gly 190	Ser	
50								٠								
	(2)	INFO	ORMA	NOI	FOR	SEQ	ID i	NO: 9	558:							

(i) SEQUENCE CHARACTERISTICS:

55

(A) LENGTH: 38 amino acids

- (B) TYPE: amino acid(D) TOPOLOGY: linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 658:
- 60 Glu Asp Asp Gly Phe Asn Arg Ser Ile His Glu Val Ile Leu Lys Asn

	1				5					10					15	
5	Ile	Thr	Trp	Туг 20	Ser	Glu	Arg	Val	Leu 25	Thr	Glu	Ile	Ser	Leu 30	Gly	Seŗ
	Leu	Leu	Ile 35	Leu	Val	Val										
10	(2)	INF	ORMA!	rion	FOR	SEQ	ֹם מוֹ	NO: (	559:							
			(i)	SEQU.	ENCE	CHA	RACT	ERIS	TICS	:						
15			(xi)	(	A) L B) T D) T UENC	YPE: OPOL	ami OGY:	no a lin	cid ear			: 65	9:			
	λ×α	Ψh∽	Ile	Cln	<b>™</b> ~~	) cn	Mob		7~~	ωρ∼	N 20000	3.00	T		<b>.</b>	***2 -
20	1	1111,	116	GIII	5	ASII	Mec	1111	AIG	10	Arg	ASD	гàг	TYE.	15	HIS
	Thr	Asn	Cys	Leu 20	Ala	Ala	Leu	Ala	Asn 25	Met	Ser	Ala	Gln	Phe 30	Arg	Ser
25	Leu	His	Gln 35	Tyr	Ala	Ala	Gln	Arg 40	Ile	Ile	Ser	Leu	Phe 45	Ser	Leu	Leu
٠	Ser	Lys 50	Lys	His	Asn										•	
30																
			•													
	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: 6	560:							
35			(i)	(	ENCE A) L B) T D) T	engt Ype:	H: 5 ami	6 am no a	ino cid		s					
40			(xi)	•	•					EQ I	ON C	: 66	0:			
40	Ser 1	Cys	Leu	Thr	Asn 5	Ser	Leu	His	His	Asn 10	Pro	Asn	Leu	Val	Tyr 15	Ala
45	Leu	Leu	Tyr	Lys 20		Asp	Leu	Phe	Glu 25	Gln	Phe	Arg	Thr	His 30	Pro	Ser
	Phe	Gln	Asp 35	Ile	Met	Gln	.Asn	Ile 40	Asp	Leu	Val	Ile	Ser 45	Phe	Phe	Ser
50	Ser	Arg 50	Leu	Leu	Gln	Ala	Gly 55	Ser								
55	(2)		ORMAT											-		

(A) LENGTH: 31 amino acids(B) TYPE: amino acid

(D) TOPOLOGY: linear

			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N:S	EQ I	D NO	: 66	1:			
5	Lys 1	Lys	His	Asn	Lys 5	Val	Leu	Glu	Gln	Ala 10	Thr	Gln	Ser	Leu	Arg 15	Gly
•	Ser	Leu	Ser	Ser 20	Asn	Asp	Val	Pro	Leu 25	Pro	Asp	Tyr	Ala	Gln 30	Asp	
10	(2)	INF	ORMA	NOIT	FOR	SEQ	ID I	NO: (	562:							
15	-		(i) :	(	A) L B) T D) T	ENGT YPE: OPOL	H: 1 ami OGY:	25 a no a lin	mino cid ear	aci		: 66	2:`			
20	Met 1	Ala	Asp	Ile	Gln 5	Thr	Glu	Arg	Ala	Tyr 10	Gln	Lys	Gln	Pro	Thr 15	Ile
	Phe	Gln	Asn	Lys 20	Lys	Arg	Val	Leu	Leu 25	Gly	Glu	Thr	Gly	Lys 30	Glu	Lys
25	Leu	Pro	Arg 35	Val	Thr	Asn	Lys	Asn 40	Ile	Gly	Leu	Gly	Phe 45	Lys	Asp	Thr
30	Pro	Arg 50	Arg	Leu	Leu	Arg	Gly 55	Thr	Tyr	Ile	Asp	Lys 60	Lys	Cys	Pro	Phe
	Thr 65	Gly	Asn	Val	Ser	Ile 70		Gly	Arg	Ile	Leu 75	Ser	Gly	Val	Val	Thr 80
35	Gln	Asp	Glu	Asp	Ala 85	Glu	Asp	His	Cys	His 90	Pro	Pro	Arg	Leu	Ser 95	Ala
	Leu	His	Pro	Gln 100	Val	Gln	Pro	Leu	Arg 105	Glu	Ala	Pro	Gln	Glu 110	His	Val
40	Cys	Thr	Pro 115	Val	Pro	Leu	Leu	Gln 120	Gly	Arg	Pro	Asp	Arg 125			
45	(2)		ORMAI							ŧ				٠		
50			(xi)	() ()	B) T	YPE: OPOL	H: 7 ami: OGY: SCRII	no a	cid ear			: 66:	3:			
55	Met 1	Lys	Met	Gln	Arg 5	Thr	Ile	Val	Ile	Arg 10	Arg	Asp	Tyr	Leu	His 15	Tyr
	Ile	Arg	Lys	Tyr 20	Asn	Arg	Phe	Glu	Lys 25	Arg	His	Lys	Asn	Met 30	Ser	Val

His Leu Ser Pro Cys Phe Arg Asp Val Gln Ile Gly Asp Ile Val Thr 35 40 45

WO 98/54963 PCT/US98/11422

	Val	GLy 50	Glu	Cys	Arg	?±'5	Leu 55	Ser	Lys	722	Val	eo Yrg	Phe	Ast.	vai	Le:	
5	Lys 65	Vai	Thr	<u>ly</u> s	Ala	Ala 70	Gly	Thr	Lys	Lys	Gln 75	Phe	31	Lys	Ζė		
10	(2)	<u> ಬ</u> ಡ್	OPMA:	CION	FOR	ಹಾರ	ID 1	10: <del>(</del>	564:								
15				(	A) L B) T D) T	eng: YPE: OPOL	H: 3 ami CGY:	0 am no a lim	ino : cii ear	acid		: 65-	<b>:</b> :				
20	Met 1	Ala	Asp	Ile	Gln 5	Thr	Glu	Arş	Ala	क्ति≆ 10	Gln	Lys	3lin	?≃:	∓== 13	Ile	
	Phe	Gln	Asn	≟уs 20 `	Lys	Arg	Val	Le:	<b>Leu</b> 25	Gly	Glu	Thr	Sly	Lys 30			
25	(2)	೨೯೮	OFMA!	TON	FCR	SEQ	ID I	<b>70:</b> :	565:								
30				(	A) L B) T D) T	engi Ype: Opol	H: 5 a <del>ri</del> CGY:	8 am no a lim	iro cid ear	acid		: 66	Ē:				
35	Lys 1	Leu	Pro	æg	∵al 5	Thr	Ast.	L; s	Asn	::e ::	Gly	leu	Gly	Phe	173 13	æp	
	Thr	Pro	Æg	Arg 20	Leu	Leu	Atg	Gl:	Ter 35	Tyr	Ile	qek	Lys	Lys 31	ಯಿತ	Pro	
40	Phe	Thr	Gly 35	Asn	7al	Ser	Ile	Arg 41	Gly	æg	Ile	Leu	Ser 45	31.7	:: <u>:</u>	Val	
45	Thr	G <u>ln</u> 50	Asp	Glu	Asp	Ala	Glu 55	Asp	æs	C):s							
50	(2)			TION SEQU		_											
30			(-)	(	A) L B) I	ENG. YPE:		8 am	eino cid		s						
55			(xi)	SEQ	UENC	E DE	SCRI	PTIC	N: S	Bį I	ON C	: 66	5:				
	His 1		His	Pro	?ro 5		Leu	Ser	Ala	Leu 10	Зis	Pro	G <u>1-</u>	∵al	3in 15	Pro	
6O	Leu	Arg	Glu	Ala	Pro	Gln	Glu	His	Val	Cys	Thr	220	Val	225	نايمة	Leu	

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Gln Gly Arg Pro Asp Arg
 5
      (2) INFORMATION FOR SEQ ID NO: 667:
             (i) SEQUENCE CHARACTERISTICS:
10
                    (A) LENGTH: 36 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 667:
15
     Met Lys Met Gln Arg Thr Ile Val Ile Arg Arg Asp Tyr Leu His Tyr
                       5
                                          10
      Ile Arg Lys Tyr Asn Arg Phe Glu Lys Arg His Lys Asn Met Ser Val
                             . 25
20
      His Leu Ser Pro
              35
25
      (2) INFORMATION FOR SEQ ID NO: 668:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 43 amino acids
30
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 668:
      Cys Phe Arg Asp Val Gln Ile Gly Asp Ile Val Thr Val Gly Glu Cys
35
      Arg Pro Leu Ser Lys Thr Val Arg Phe Asn Val Leu Lys Val Thr Lys
                   20
                                      25
40
      Ala Ala Gly Thr Lys Lys Gln Phe Gln Lys Phe
              35
                   .
45
      (2) INFORMATION FOR SEQ ID NO: 669:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 33 amino acids
                    (B) TYPE: amino acid
50
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 669:
      Pro Arg Arg Leu Leu Arg Gly Thr Tyr Ile Asp Lys Lys Cys Pro Phe
55
      Thr Gly Asn Val Ser Ile Arg Gly Arg Ile Leu Ser Gly Val Val Thr
```

Gln

5	(2) INFORMATION FOR SEQ ID NO: 670:
3	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 60 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 670:
	Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met 1 5 10 15
15	Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg 20 25 30
20	Leu Ile Cys Lys Gly Thr Ile Glu Glu Arg Ile Leu Gln Arg Ala Lys 35 40 45
20	Glu Lys Ser Glu Ile Gln Arg Met Val Ile Ser Gly 50 55 60
25	(2) INFORMATION FOR SEQ ID NO: 671:
30	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 67 amino acids  (B) TYPE: amino acid
	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 671:
35	Thr Arg Met Ile Asp Leu Leu Glu Glu Tyr Met Val Tyr Arg Lys His  1 5 10 15
	Thr Tyr Xaa Arg Leu Asp Gly Ser Ser Lys Ile Ser Glu Arg Arg Asp 20 25 30
40	Met Val Ala Asp Phe Gln Asn Arg Asn Asp Ile Phe Val Phe Leu Leu 35 40 45
45	Ser Thr Arg Ala Gly Gly Leu Gly Ile Asn Leu Thr Ala Xaa Asp Thr 50 55 60
,,,	Val His Phe 65
50	(2) INFORMATION FOR SEQ ID NO: 672:
55	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 32 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 672:
60	Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met  1 5 10 15

PCT/US98/11422

Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg 25 5 10 (2) INFORMATION FOR SEQ ID NO: 673: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 31 amino acids (B) TYPE: amino acid 15 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 673: Val Tyr Arg Leu Ile Cys Lys Gly Thr Ile Glu Glu Arg Ile Leu Gln 20 Arg Ala Lys Glu Lys Ser Glu Ile Gln Arg Met Val Ile Ser Gly 25 25 (2) INFORMATION FOR SEQ ID NO: 674: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 33 amino acids 30 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 674: Thr Arg Met Ile Asp Leu Leu Glu Glu Tyr Met Val Tyr Arg Lys His 35 Thr Tyr Xaa Arg Leu Asp Gly Ser Ser Lys Ile Ser Glu Arg Arg Asp 20 25 40 Met 45 (2) INFORMATION FOR SEQ ID NO: 675: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 38 amino acids (B) TYPE: amino acid 50 (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 675: Arg Arg Asp Met Val Ala Asp Phe Gln Asn Arg Asn Asp Ile Phe Val 5 55 Phe Leu Leu Ser Thr Arg Ala Gly Gly Leu Gly Ile Asn Leu Thr Ala 20 25 Xaa Asp Thr Val His Phe 60 35

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(2) INFORMATION FOR SEQ ID NO: 676:
 5
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 37 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
10
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 676:
      Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met
15
     Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg
                                      25
      Leu Ile Cys Lys Gly
              35
20
      (2) INFORMATION FOR SEQ ID NO: 677:
25
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 37 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 677:
30
      Ile Phe Tyr Asp Ser Asp Trp Asn Pro Thr Val Asp Gln Gln Ala Met
                        5
       1
      Asp Arg Ala His Arg Leu Gly Gln Thr Lys Gln Val Thr Val Tyr Arg
35
                                       25
      Leu Ile Cys Lys Gly
               35
40
      (2) INFORMATION FOR SEQ ID NO: 678:
             (i) SEQUENCE CHARACTERISTICS:
45
                    (A) LENGTH: 29 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 678:
50
      Arg Leu Ile Cys Lys Gly Thr Ile Glu Glu Arg Ile Leu Gln Arg Ala
                                           10
      Lys Glu Lys Ser Glu Ile Gln Arg Met Val Ile Ser Gly
                                       25
55
      (2) INFORMATION FOR SEQ ID NO: 679:
60
            (i) SEQUENCE CHARACTERISTICS:
```

				(	B) T	ENGT YPE: OPOL	ami	no a	cid	aci	ds					
5			(xi)	-						EQ I	D NO	: 67	9:			
J	Met 1	Ser	Leu	His	Gly 5	Lys	Arg	Lys	Glu	Ile 10	Tyr	Lys	тут	Glu	Ala 15	Pro
10	Trp	Thr	Val	Туг 20	Ala	Met	Asn	Trp	Ser 25	Val	Arg	Pro	Asp	Lys 30	Arg	Phe
	Arg	Leu	Ala 35	Leu	Gly	Ser	Phe	Val 40	Glu	Glu	Tyr	Asn	Asn 45	-	Val	Ğln
15	Leu	Val 50	Gly	Leu	Asp	Glu	Glu 55	Ser	Ser	Glu	Phe	Ile 60	Cys	Arg	Asn	Thr
20	Phe 65	Asp	His	Pro	Туг	Pro 70	Thr	Thr	Lys	Leu	Met 75	Trp	Ile	Pro	Asp	Thr 80
	Lys	Gly	Val	Tyr	Pro 85	Asp	Leu	Leu	Ala	Thr 90	Ser	Gly	Asp	Tyr	Leu 95	Arg
25	Val	Trp	Arg	Val 100	Gly	Glu	Thr	Glu	Thr 105	Arg	Leu	Glu	Cys	Leu 110	Leu	Asn
	Asn	Asn	Lys 115	Asn	.Ser	qzA	Phe	Cys 120	Ala	Pro	Leu	Thr	Ser 125	Phe	Asp	Trp
30	Asn	Glu 130	Val	Asp	Pro	Tyr	Leu 135	Leu	Gly	Thr	Ser	Ser 140	Ile	Asp	Thr	Thr
35	Cys 145	Thr	Ile	Trp	Gly	Leu 150	Glu	Thr	Gly	Gln	Val 155	Leu	Gly	Arg	Val	Asn 160
	Leu	Val	Ser	Gly	His 165	Val	Lys	Thr	Gln	Leu 170	Ile	Ala	His	Asp	Lys 175	Glu
40	Val	Tyr	Asp	Ile 180	Ala	Phe	Ser	Arg	Ala 185	Gly	Gly	Gly	Arg	Asp 190	Met	Phe
	Ala	Ser	Val 195	Gly	Ala	Asp	Gly	Ser 200	Val	Arg	Met	Phe	Asp 205	Leu	Arg	His
45	Leu	Glu 210	His	Ser	Thr	Ile	Ile 215	Tyr	Glu	Asp	Pro	Gln 220	His	His	Pro	Leu
50	Leu 225	Arg	Leu	Cys	Trp	Asn 230	Lys	Gln	Asp	Pro	Asn 235	Tyr	Leu	Ala	Ţhr	Met 240
50	Ala	Met	Asp	Gly	Met 245	Glu	Val	Val	Ile	Leu 250	Asp	Val	Arg	Val	Pro 255	Ala
55	His	Leu	Xaa	Pro 260	Gly	Thr	Thr	Ile	Glu 265	His	Val	Ser	Met	Ala 270	Leu	Leu
	Gly	Pro	His 275	Ile	His	Pro	Ala	Thr 280	Ser	Ala	Leu	Gln	Arg 285	Met	Thr	Thr
60	Arg	Leu	Ser	Ser	Gly	Thr	Ser	Ser	Lys	Cys	Pro	Glu	Pro	Leu	Arg	Thr

PCT/US98/11422

		290					295					300				
5	Leu 305	Ser	Trp	Pro	Thr	Gln 310	Leu	Xaa	Gly	Glu	Ile 315	Asn	Asn	Val	Gln	Trp 320
3	Ala	Ser	Thr	Gln	Pro 325	Glu	Leu	Ser	Pro	Ser 330		Thr	Thr	Thr	Ala 335	Trp
10	Arg	Tyr	Ser	Glu 340	Cys	Ser	Val	Gly	Gly 345	Ala	Val	Pro	Thr	Arg 350	Gln	Gly
•	Leu	Leu	Тут 355	Phe	Leu	Pro	Leu	Pro 360	His	Pro	Gln	Ser				
15										٠						
	(2)	INF	ORMA	NOI	FOR	SEQ	ID i	vo: (	680:						•	
20			(i) :	. (	A) L B) T D) T	CHAI ENGT: YPE: OPOL E DE:	H: 1 ami OGY:	36 a no a lin	mino cid ear	aci		: 68	0 :			
25	Met 1	Ser	Leu	His	Gly 5	Lys	Arg	Lys	Glu	Ile 10	Tyr	Lys	Tyr	Glu	Ala 15	Pro
30	Trp	Thr	Val	Tyr 20	Ala	Met	Asn	Trp	Ser 25	Val	Arg	Pro	Asp	Lys 30	Arg	Phe
30	Arg	Leu	Ala 35	Leu	Gly	Ser	Phe	Val 40	Glu	Glu	Tyr	Asn	Asn 45	Lys	Val	Gln
35	Leu	Val 50	_	Leu	Asp	Glu	Glu 55	Ser	Ser	Glu	Phe	Ile 60	Суѕ	Arg	Asn	Thr
	Phe 65	Asp	His	Pro	Tyr	Pro 70	Tbr	Thr	Lys	Leu	Met 75	Trp	Ile	Pro	Asp	Thr 80
40	Lys	Gly	Val	Tyr	Pro 85	Asp	Leu	Leu	Ala	Thr 90	Ser	Gly	Asp	Tyr	Leu 95	Arg
45	Val	Trp	Arg	Val 100	Gly	Glu	Thr	Glu	Thr 105	Arg	Leu	Glu	Cys	Leu 110	Leu	Asn
	Asn	Asn	Lys 115	Asn	Ser	Asp	Phe	Cys 120	Ala	Pro	Leu	Thr	Ser 125	Phe	Asp	Trp
50	Asn	Glu 130	.Val	Asp	Pro	Tyr	Leu 135	Leu								
55	(2)	INF	ORMA	SEQU (	ENCE A) L B) T	SEQ CHA ENGT YPE:	RACT H: 1 ami	ERIS 40 a no a	TICS mino		ds.				·	
60			(xi)			E DE				EQ I	D NO	: 68	1:			

	Ser 1	Phe	Asp	Trp	Asn 5	Glu	Val	Asp	Pro	Tyr 10	Leu	Leu	Gly	Thr	Ser 15	Ser
5	Ile	Asp	Thr	Thr 20	Cys	Thr	Ile	Trp	Gly 25	Leu	Glu	Thr	Gly	Gln 30	Val	Leu
10	Gly	Arg	Val 35	Asn	Leu	Val	Ser	Gly 40	His	Val	Lys	Thr	Gln 45	Leu	Ile	Ala
	His	Asp 50	Lys	Glu	Val	Tyr	Asp 55	Ile	Ala	Phe	Ser	Arg 60	Ala	Gly	Gly	Gly
15	Arg 65	Asp	Met	Phe	Ala	Ser 70	Val	Gly	Ala	Asp	Gly 75	Ser	Val	Arg	Met	Phe 80
	Asp	Leu	Arg	His	Leu 85	Glu	His	Ser	Thr	Ile 90	Ile	Tyr	Glu	Asp	Pro 95	Gln
20	His	His	Pro	Leu 100	Leu	Arg	Leu	Cys	Trp 105	Asn	Lys	Gln	Asp	Pro 110	Asn	Tyr
25	Leu	Ala	Thr 115	Met	Ala	Met	Asp	Gly 120	Met <sub>.</sub>	Glu	Val	Val	Ile 125	Leu	Asp	Val
	Arg	Val 130	Pro	Ala	His	Leu	Хаа 135	Pro	Gly	Thr	Thr	Ile 140				
30	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	VO: 6	82:						•	
35			(i) :	(	A) L B) T	ENGT YPE :	RACTI H: 1 ami OGY:	70 a no a	mino cid		ds					
			(xi)	SEQ						EQ I	D NO	: 68	2:			
40	Val 1	Gly	Ala	Asp	Gly 5	Ser	Val	Arg	Met	Phe 10	Asp	Leu	Arg	His	Leu 15	Glu
	His	Ser	Thr	Ile 20	Ile	Tyr	Glu	Asp	Pro 25	Gln	His	His	Pro	Leu 30	Leu	Arg
45	Leu	Cys	Trp 35	Asn	Lys	Gln	Asp	Pro 40	Asn	Tyr	Leu	Ala	Thr 45	Met	Ala	Met
50	Asp	Gly 50	Met	Glu	Val	Val	Ile 55	Leu	Asp	Val	Arg	Val 60	Pro	Ala	His	Leu
	Xaa 65	Pro	Gly	Thr	Thr	Ile 70	Glu	His	Val	Ser	Met 75	Ala	Leu	Leu	Gly	Pro 80
55	His	Ile	His	Pro	Ala 85	Thr	Ser	Ala	Lėu	Gln 90	Arg	Met	Thr	Thr	Arg 95	Leu
	Ser	Ser	Gly	Thr	Ser	Ser	Lys	Cys	Pro	Glu	Pro	Leu	Arg	Thr	Leu	Ser
				100					105					110		•

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			115					120					125			
·5	Thr	Gln 130	Pro	Glu	Leu	Ser	Pro 135	Ser	Ala	Thr	Thr	Thr 140	Ala	Trp	Arg	Tyr
J	Ser 145	Glu	Cys	Ser	Val	Gly 150	Gly	Ala	Val	Pro	Thr 155	Arg	Gln	Gly	Leu	Leu 160
10	Tyr	Phe	Leu	Pro	Leu 165	Pro	His	Pro	Gln	Ser 170						
15	(2)	INF		rion												
70				(	A) L B) T D) T	ENGT YPE : OPOL	H: 2 ami OGY:	86 a no a lin	mino cid ear	aci						
20			(xi)	SEQ	UENC:	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 68	3:			
	Leu 1	Tyr	Ala	Thr	Ala 5	Thr	Val	Ile	Ser	Ser 10	Pro	Ser	Thr	Glu	Хаа 15	Leu
25	Ser	Gln	Asp	Gln 20	Gly	Asp	Arg	Ala	Ser 25	Leu	Asp	Ala	Ala	Asp 30	Ser	Gly
30	Arg	Gly	Ser 35	Trp	Thr	Ser	Cys	Ser 40	Ser	Gly	Ser	His	Asp 45	Asn	Ile	Gln
,	Thr	Ile 50	Gln	His	Gln	Arg	Ser 55	Trp	Glu	Thr	Leu	Pro 60	Phe	Gly	His	Thr
35	His 65	Phe	Asp	Tyr	Ser	Gly 70	Asp	Pro	Ala	Gly	Leu 75	Trp	Ala	Ser	Ser	Ser 80
	His	Met	Asp	Gln	Ile 85	Met	Phe	Ser	Asp	His 90	Ser	Thr	Lys	Tyr	Asn 95	Arg
<b>4</b> 0	Gln	Asn	Gln	Ser 100	Arg	Glu	Ser	Leu	Glu 105	Gln	Ala	Gln	Ser	Arg 110	Ala	Ser
15	Trp	Ala	Ser 115	Ser	Thr	Gly	Tyr	Trp 120	Gly	Glu	Asp	Ser	Glu 125	Gly	Asp	Thr
	Gly	Thr 130	Ile	Lys	Arg	Arg	Gly 135	Gly	Lys	Asp	Val	Ser 140	Ile	Glu	Ala	Glu
50	Ser 145	Ser	Ser	Leu	Thr	Ser 150	Val	Thr	Thr	Glu	Glu 155	Thr	Lys	Pro	Val	Pro 160
	Met	Pro	Ala	His	Ile 165	Ala	Val	Ala	Ser	Ser 170	Thr	Thr	Lys	Gly	Leu 175	Ile
55	Ala	Arg	Lys <sub>.</sub>	Glu 180	Gly	Arg	Tyr	Arg	Glu 185	Pro	Pro	Pro	Thr	Pro 190	Pro	Gly
50	Tyr	Ile	Gly 195	Ile	Pro	Ile	Thr	Asp 200	Phe	Pro	Glu	Gly	His 205	Ser	His	Pro

	Ala	Arg 210	Lys	Pro	Pro	Asp	Tyr 215	Asn	Val	Ala	Leu	Gln 220	Arg	Ser	Arg	Met
5	Val 225	Ala	Arg	Ser	Ser	Asp 230	Thr	Ala	Gly	Pro	Ser 235	Ser	Val	Gln	Gln	Pro 240
	His	Gly	His	Pro	Thr 245	Ser	Ser	Ärg	Pro	Val 250	Asn	Lys	Pro	Gln	Trp 255	His
10	Lys	Xaa	Asn	Glu 260	Ser	Asp	Pro	Arg	Leu 265	Ala	Pro	Туг	Gln	Ser 270	Gln	Gly
15	Phe	Ser	Thr 275	Glu ,	Glu	Asp	Glu	Asp 280	Glu	Gln	Val	Ser	Ala 285	Val		
	(2)	INFO	ORMAT	rion	FOR	SEQ	ID 1	NO: 6	84:							
20			(i) :	(:	ENCE A) L B) T D) T	ENGT YPE:	H: 4	2 am no a	ino a		s					
25			(xi)	SEQ	JENC!	E DE	SCRI	PTIO	N: SI	EQ II	ОИС	: 68	4:			
	His 1	Met	Asp	Gln	Ile 5	Met	Phe	Ser	Asp	His 10	Ser	Thr	Lys	Tyr	Asn 15	Arg
30	,Gln	Asn	Gln	Ser 20	Arg	Glu	Ser	Leu	Glu 25	Gln	Ala	Gln	Ser	Arg 30	Ala	Ser
	Trp	Ala	Ser 35	Ser	Thr	Gly	Tyr	Trp 40	Gly	Glu						
35																
	(2)	INF	ORMA!	rion	FOR	SEQ	IQ.	NO: 6	585:							
40				(	A) L B) T D) T	ENGT YPE: OPOL	H: 5 ami OGY:	l am no a lin	ino d cid ear	acid		: 68:	5:			
45	Ser 1	Val	Thr	Thr	Glu 5	Glu	Thr	Lys	Pro	Val 10	Pro	Met	Pro	Ala	His 15	
50	Ala	Val	Ala	Ser 20	Ser	Thr	Thr	Lys	Gly 25	Leu	Ile	Ala	Arg	Lys 30	Glu	Gly
50	Arg	Tyr	Arg 35	Glu	Pro	Pro	Pro	Thr 40	Pro	Pro	Gly	Tyr	Ile 45	Gly	Ile	Pro
55	Ile	Thr 50	Asp									•				

(2) INFORMATION FOR SEQ ID NO: 686:

	(i) SEQUENCE CHREACTERISTICS:  (A) LENGTH: 57 amino acids  (B) TYPE: amino acid	
	(D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 686:	
	Val Ala Leu Glm Arg Ser Arg Met Val Ala Arg Ser Ser Asp Thr Al 1 5 10 15	a
10	Gly Pro Ser Ser Val Glm Glm Pro His Gly His Pro Thr Ser Ser Ar 20 25 30	g
15	Pro Val Asn Lys Pro Gln Trp His Lys Xaa Asn Glu Ser Asp Pro Ar 35 40 45	g.
13	Leu Ala Pro Tyr Gln Ser Gln Gly Phe 50 55	
20	(2) DIFORMATION FOR SEQ ID NO: 687:	
25	(i) SEQUENCE CHAPACTERISTICS:  (A) LENGTH: 41 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 687:	
30	Cys Leu Leu Phe Val Phe Val Ser Leu Gly Met Arg Cys Leu Phe Tr 1 5 10 15	p
	Thr Ile Val Tyr Asn Val Leu Tyr Leu Lys His Lys Cys Asn Thr Va	ıı
35	Leu Leu Cys Tyr His Leu Cys Ser Ile 35 40	
40	(2) INFORMATION FOR SEQ ID NC: 688:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 67 amino acids  (B) TYPE: amino acid	
45	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 688:	
50	Ala Cys Ser Lys Leu Ile Pro Ala Phe Glu Met Val Met Arg Ala Ly 1 5 10 15	'S
50	Asp Asn Val Tyr His Leu Asp Cys Phe Ala Cys Gln Leu Cys Asn Gl 20 25 30	n.
55	Arg Xaa Cys Val Gly Asp Lys Phe Phe Leu Lys Asn Asn Xaa Xaa Le 35 40 45	u
	Cys Gln Thr Asp Tyr Glu Glu Gly Leu Met Lys Glu Gly Tyr Ala Pr 50 55 60	0
60	Xaa Val Arg	

	•	
5	(2) INFORMATION FOR SEQ ID NO: 689:	
10	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 45 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 689:</li> </ul>	
	Ser Ala Leu Ser Glu Pro Gly Ala Pro Asp Arg Arg Pro Cys Pro  1 5 10 15	
15		
	Glu Ser Val Pro Arg Arg Pro Asp Asp Glu Gln Trp Pro Pro Pro Thr 20 25 30	
20	Ala Leu Cys Leu Asp Val Ala Pro Leu Pro Pro Ser Ser 35 40 45	
25	(2) INFORMATION FOR SEQ ID NO: 690:	
23	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 43 amino acids (B) TYPE: amino acid	
30	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 690:	
	Pro Val Gly Tyr Leu Asp Lys Gln Val Pro Asp Thr Ser Val Gln Glu	
25	1 5 10 15	
35	Thr Asp Arg Ile Leu Val Glu Lys Arg Cys Trp Asp Ile Ala Leu Gly 20 25 30	
	Pro Leu Lys Gln Ile Pro Met Asn Leu Phe Ile 35	
40		
	(2) INFORMATION FOR SEQ ID NO: 691:	
45	(i) SEQUENCE CHARACTERISTICS:	
	(A) LENGTH: 214 amino acids (B) TYPE: amino acid	
50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 691:	
	Ala His Ala Ser Glu Ser Gly Glu Arg Trp Trp Ala Cys Cys Gly Val 1 5 10 15	
<i></i>	Arg Phe Gly Leu Arg Ser Ile Glu Ala Ile Gly Arg Ser Cys Cys His	
55	20 25 30	
	Asp Gly Pro Gly Gly Leu Val Ala Asn Arg Gly Arg Arg Phe Lys Trp 35 40 45	
60	Ala Ile Glu Leu Ser Gly Pro Gly Gly Gly Ser Arg Gly Arg Ser Asp	

		50					55					60				
5	Arg 65		Ser	Gly	Gln	Gly 70	Asp	Ser	Leu	Tyr	Pro 75	Val	Gly	Tyr	Leu	Asp 08
J	Lys	Gln	Val	Pro	Asp 85	Thr	Ser	Val	Gln	Glu 90	Thr	Asp	Arg	Ile	Leu 95	Val
10	Glu	Lys	Arg	Суs 100	Trp	Asp	Ile	Ala	Leu 105	Gly	Pro	Leu	Lys	Gln 110	Ile	Pro
	Met	Asn	Leu 115	Phe	Ile	Met	Тут	Met 120	Ala	Gly	Asn	Thr	Ile 125	Ser	Ile	Phe
15	Pro	Thr 130	Met	Met	Val	Суз	Met 135		Ala	Trp	Arg	Pro 140	Ile	Gln	Ala	Leu
20	Met 145	Ala	Ile	Ser	Ala	Thr 150	Phe	Lys	Met	Leu	Glu 155	Ser	Ser	Ser	Gln	Lys 160
	Phe	Leu	Gln	Gly	Leu 165	Val	Tyr	Leu	Ile	Gly 170	Asn	Leu	Met	Gly	Leu 175	Ala
25	Leu	Ala	Val	Туг 180	Lys	Cys	Gln	Ser	Met 185	Gly	Leu	Leu	Pro	Thr 190	His	Äla
	Ser	Asp	Trp 195	Leu	Ala	Phe	Ile	Glu 200	Pro	Pro	Glu	Arg	Met 205	Glu	Phe	Ser
30	Gly	Gly 210	Gly	Leu	Leu	Leu								•		
35	(2)	INFO	ORMA'	NOIT	FOR	SEQ	ID I	NO: 6	592:							
40				(	A) L B) T D) T	ENGT YPE: OPOL	H: 4 ami OGY:	6 am no a lin	ino cid ear	acid		: 69:	2:			
45	Ala 1	Thr	Phe	Lys	Met 5	Leu	Glu	Ser	Ser	Ser 10	Gln	Lys	Phe	Leu	Gln 15	Gly
43	Leu	Val	Tyr	Leu 20	Ile	Gly	Asn	Leu	Met 25	Gly	Leu	Ala	Leu	Ala 30	Val	Туг
50	Lys	Суз	Gln 35	Ser	Met	Gly	Leu	Leu 40	Pro	Thr	His	Ala	Ser 45	Asp	٠	
55	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	NO: 6	593 :							
			(i)		ENCE A) L B) T	ENGT	H: 4	3 am	ino		s					
60				(	D) T	OPOL	OGY:	lin	ear				_			

	Pro 1		Gly	Tyr	Leu 5	Asp	Lys	Gln	Val	Pro 10	Asp	Thr	Ser	Val	Gln 15	Glu
5	Thr	Asp	Arg	Ile 20	Leu	Val	Glu	Lys	Arg -25	Cys	Trp	Asp	Ile	Ala 30	Leu	Gly
10	Pro	Leu	Lys 35	Gln	Ile	Pro	Met	Asn 40	Leu	Phe	Ile					
15	(2)	INF	ORMA:	SEQUI ( ) (	ENCE A) L B) T D) T	CHA ENGT YPE: OPOL	RACT H: 4 ami OGY:	ERIS 8 am no a lin	TICS ino cid ear	acid		· 69	<b>4</b> •			
20	Pro 1	Thr												Ile	Gln 15	Ile
25	Arg	Phe	Pro	Ser 20	Phe	Tyr	His	Lys	Leu 25	Val	Asp	Ser	Gly	Arg 30	Met	Arg
20	Ser	Lys	Arg 35	Glu	Thr	Arg	Arg	Glu 40	Asp	Ser	Asp	Thr	Lys 45	His	Asn	Leu
30																
35	(2)	INFO														
40			(i) :	(. ()	A) L B) T D) T	ENGT YPE: OPOL	H: 1 ami OGY:	67 a no a lin	mino cid ear	aci		: 69	5:			
45	Thr 1	Glu	His	Ile	Ile 5	Ala	Val	Met	Ile	Thr 10	Glu	Leu	Arg	Gly	Lys 15	Asp
	Ile	Leu	Ser	Туг 20	Leu	Glu	Lys	Asn	Ile 25	Ser	Val	Gln	Met	Thr 30	Ile	Ala
50	Val	Gly	Thr 35	Arg	Met	Pro	Pro	Lys 40	Asn	Phe	Ser	Arg	Gly 45	Ser	Leu	Val
	Phe	Val 50	Ser	Ile	Ser	Phe	Ile 55	Val	Leu	Met	Ile	Ile 60	Ser	Ser	Ala	Trp
55	Leu 65		Phe	Tyr	Phe	Ile 70	Gln	Lys	Ile	Arg	Tyr 75	Thr	Asn	Ala	Arg	Asp 80
60	Arg	Asn	Gln	Arg	Arg 85	Leu	Gly	Asp	Ala	Ala 90	Lys	Lys	Ala	Ile	Ser 95	Lys

	Leu	Thr	Thr	Arg 100	Thr	Val	Lys	Lys	Gly 105	Asp	Lys	Glu		Asp 110	Pro	Asp
5	Phe	Asp	His 115	Суз	Ala	Val	Cys	Ile 120	Glu	Ser	Tyr	Lys	Gln 125	Asn	Asp	Val
	Val	Arg 130	Ile	Leu	Pro	Cys	Lys 135	His	Val	Phe	His	Lys 140	Ser	Cys	Val	Asp
10	Pro 145	Trp	Leu	Ser	Glu	His 150	Cys	Thr	Cys	Pro	Met 155	Суз	Lys	Leu	Asn	Ile 160
15	Leu	Lys	Ala	Leu	Gly 165	Ile	Val									
	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	NO: 6	596:							
20				(	A) L B) T D) T	ENGT YPE: OPOL	H: 2 ami OGY:	76 a no a lin	mino cid ear	aci			٠.			
25	15-4-									EQ I					,	
	met 1	Thr	HIS	Pro	GIY GIY	Thr	GIu	His	Ile	Ile 10	Ala	Val	Met	Ile	Thr 15	Glu
30	Leu	Arg	Gly	Lys 20	Asp	Ile	Leu	Ser	Туг 25	Leu	Glu	Lys	Asn	Ile 30	Ser	Val
	Gln	Met	Thr 35	Ile	Ala	Val	Gly	Thr 40	Arg	Met	Pro	Pro	Lys 45	Asn	Phe	Ser
35	Arg	Gly 50	Ser	Leu	Val	Phe	Val 55	Ser	Ile	Ser	Phe	Ile 60	Val	Leu	Met	Ile
40	Ile 65	Ser	Ser	Ala	Trp	Leu 70	Ile	Phe	Tyr	Phe	Ile 75	Gln	Lys	Ile	Arg	Tyr 80
	Thr	Asn	Ala	Arg	Asp 85	Arg	Asn	Gln	Arg	Arg 90	Leu	Gly	Asp	Ala	Ala 95	Lys
45	Lys	Ala	Ile	Ser 100	Lys	Leu	Thr	Thr	Arg 105	Thr	Val	Lys	Lys	Gly 110	Asp	Lys
	Glu	Thr	Asp 115	Pro	Asp	Phe	Asp	His 120	Cys	Ala	Val	Суѕ	Ile 125	Glu	Ser	Tyr
50	Lys	Gln 130	Asn	Asp	Val	Val	Arg 135	Ile	Leu	Pro	Cys	Lys 140	His	Val	Phe	His
55	Lys 145	Ser	Cys	Val	Asp	Pro 150	Trp	Leu	Ser	Glu	His 155	Cys	Thr	Cys	Pro	Met 160
	Суѕ	Lys	Leu	Asn	Ile 165	Leu	Lys	Ala	Leu	Gly 170	Ile	Val	Pro	Asn	Leu 175	Pro
60	Cys	Thr	Asp	Asn	Val	Ala	Phe	Asp	Met	Glu	Arg	Leu	Thr	Arg	Thr	Gln

	Ala	Val	Asn 195	Arg	Arg	Ser	Ala	Leu 200	Gly	Asp	Leu	Ala	Gly 205	Asp	Asn	Ser
5	Leu	Gly 210	Leu	Glu	Pro	Leu	Arg 215	Thr	Ser	Gly	Ile	Ser 220	Pro	Leu	Pro	Gln
10	Asp 225	Gly	Glu	Leu	Thr	Pro 230	Arg	Thr	Gly	Glu	11e 235	Asn	Ile	Ala	Val	Thr 240
	Lys	Glu	Trp	Phe	Ile 245	Ile	Ala	Ser	Phe	Gly 250	Leu	Leu	Ser	Ala	Leu 255	Thr
15	Leu	Cys	Tyr	Met 260	Ile	Ile	Arg	Ala	Thr 265	Ala	Ser	Leu	Asn	Ala 270	Asn	Glu
	Val	Glu	Trp 275	Phe												
20					• ,											
	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	<b>10:</b> (	597 :							
25				(	A) L B) T D) T	ENGT YPE : OPOL	H: 6 ami OGY:	9 am no a lin	ino cid ear	acid		• 69'	7 ·			
30	ጥኮኍ			Ile						_				C111	Tare	) an
	1	Giu	1112		5	ALG	Vai	Mec	116	10	Giu	Ten	My	GIY	15	ASP
35	Ile	Leu	Ser	Тут 20	Leu	Glu	Lys	Asn	Ile 25	Ser	Val	Gln	Met	Thr 30	Ile	Ala
	Val	Gly	Thr 35	Arg	Met	Pro	Pro	Lys 40	Asn	Phe	Ser	Arg	Gly 45	Ser	Leu	Val
40	Phe	Val 50	Ser	Ile	Ser	Phe	Ile 55		Leu	Met	Ile	Ile 60	Ser	Ser	Ala	Trp
	Leu 65	Ile	Phe	Tyr	Phe											
45					:			•								
	(2)	INF	ORMA!	rion	FOR	SEQ	ID 1	NO: 6	698 :							
50				(	A) L B) T D) T	ENGT YPE : OPOL	H: 5 ami OGY:	8 am no a lin	ino cid ear	acid		: 69	8:			
55	Ser 1	Ile	Ser	Phe	Ile 5	Val	Leu	Met	Ile	Ile 10	Ser	Ser	Ala	Trp	Leu 15	Ile
<b>5</b> 0	Phe	Тух	Phe	Ile 20	Gln	Lys	Ile	Arg	Tyr 25	Thr	Asn	Ala	Arg	Asp 30	Arg	Asn,

	Gln	Arg	Arg 35	Leu	Gly	Asp	Ala	A1a 40	Lys	Lys	Ala	Ile	Ser 45	Lys	Leu	Thr
5	Thr	Arg 50	Thr	Val	Lys	Lys	Gly 55	Asp	Lys	Glu						
10	(2)	INFO	ORMA	rion	FOR	SEQ	ID 1	<b>1</b> 0: 6	599:				•			
			(i)	(	ENCE A) L B) T D) T	ENGT YPE:	H: 6	6 am no a	ino cid		s					
15			(xi)	SEQ	UENC	E DE	SCRI:	PTIO	N: S	EQ I	ON O	: 69	9:			
	Val 1	Lys	Lys	Gly	Asp 5	Lys	Glu	Thr	Asp	Pro 10	Asp	Phe	Asp	His	Cys 15	Ala
20	Val	Cys	Ile	Glu 20	Ser	Tyr	Lys	Gln	Asn 25	Asp	Val	Val	Arg	Ile 30	Leu	Pro
25	Cys	Lys	His 35	Val	Phe	His	Lys	Ser 40	Cys	Val	Asp	Pro	Trp 45	Leu	Ser	Glu
	His	Суs 50	Thr	Cys	Pro	Met	Cys 55	Lys	Leu	Asn	Ile	Leu 60	Lys	Ala	Leu	Gly
30	Ile 65	Val														
35	(2)	INF	ORMA'	rion	FOR	SEQ	ID 1	NO: 7	700:							
),)			(i)		ENCE A) L B) T	ENGT	н: 1	06 a	mino		ds					
10			(xi)		D) T	OPOL	OGY:	lin	ear	EQ I	D NO	: 70	0:			
	Met 1	Thr	His	Pro	Gly 5	Thr	Glu	His	Ile	Ile 10	Ala	Val	Met	Ile	Thr 15	Glu
15	Leu	Arg	Gly	Lys 20	Asp	Ile	Leu	Ser	Туr 25	Leu	Glu	Lys	Asn	Ile 30	Ser	Val
50	Gln	Met	Thr 35	Ile	Ala	Val	Gly	Thr 40	Arg	Met	Pro	Pro	Lys 45	Asn	Phe	Ser
-	Arg	Gly 50	Ser	Leu	Val	Phe	Val 55	Ser	Ile	Ser	Phe	Ile 60	Val	Leu	Met	Ile
55	Ile 65	Ser	Ser	Ala	Trp	<b>Leu</b> 70	Ile	Phe	Tyr	Phe	Ile 75	Gln	Lys	Ile	Arg	Tyr 80
	Thr	Asn	Ala	Arg	Asp 85	Arg	Asn	Gln	Arg	Arg 90	Leu	Gly	Asp	Ala	Ala 95	Lys
50	Lare	λla	Tla	Sor	Tare	T ALL	Thr	Thr	7~~	Thr						

5	(2)	INF	ORMA'	TION	FOR	SEQ	ID I	.vo: '	701:							
10		,		(	A) L B) T D) T	ENGT YPE : OPOL	H: 8 ami OGY:	4 am no a lin	ino cid ear	acid		: <b>7</b> 0.	1:			
15	Ala 1	Ala	Lys	Lys	Ala 5	Ile	Ser	Lys	Leu	Thr 10	Thr	Arg	Thr	Val	Lys 15	Lys
	Gly	Asp	Lys	Glu 20	Thr	Asp	Pro	Asp	Phe 25	Asp	His	Суѕ	Ala	Val 30	Суз	Ile
20	Glu	Ser	Tyr 35	Lys	Gl'n	Asn	Asp	Val 40	Val	Arg	Ile	Leu	Pro 45	Cys	Lys	His
	Val	Phe 50	His	Lys	Ser	Cys	Val 55	Asp	Pro	Trp	Leu	Ser 60	Glu	His	Cys	Thr
25	Cys 65	Pro	Met	Cys	Lys	Leu 70	Asn	Ile	Leu	Lys	Ala 75	Leu	Gly	Ile	Val	Pro 80
30	Asn	Leu	Pro	Cys												
	(2)	INF	ORMA	rion	FOR	SEQ	ID I	NO: '	702:							
35		•	(i)	(	A) L B) T	ENGT YPE:	H: 8 ami	6 am no a	ino cid		s					
40			(xi)	SEQ		OPOL E DE:				EQ I	D NO	: 70	2:			
	Thr 1		Ala	Val	Asn 5	Arg <sub>.</sub>	Arg	Ser	Ala	Leu 10	Gly	Asp	Leu	Ala	Gly 15	Asp
45	Asn	Ser	Leu	Gly 20	Leu	Glu	Pro	Leu	Arg 25	Thr	Ser	Gly	Ile	Ser 30	Pro	Leu
,	Pro	Gln	Asp 35	Gly	Glu	Leu	Thr	Pro 40	Arg	Thr	Gly	Glu	11e 45	Asn	Ile	Ala
50	Val	Thr 50	Lys	Gļú	Trp	Phe	Ile 55	Ile	Ala	Ser	Phe	Gly 60	Leu	Leu	Ser	Ala
55	Leu 65	Thr	Leu	Суз	Tyr	Met 70	Ile	Ile	Arg	Ala	Thr 75	Ala	Ser	Leu	Asn	Ala 80
<i></i>	Asn	Glu	Val	Glu	Trp 85	Phe										

•	2)	ΞŦ	OPYA:	TION	FCR	ಕ್ಟಾ	ID :	:T: '	703:							
5				( ( (	A) I 3) T 0) T	engi Ype : Ype :	PACT H: 3 ami OG:: SCPI	41 a mo a 'lin	mino cid ear	aci		: 70	3:			
10	Pro 1	Leu	His	Gly	7al 5	Àla	ಸಿತ್ರಾ	His	Leu	Gly 10	೦್ಯಕ	Asp	Pro	Gln	Thr 15	Arg
	Phe	Fhe	Val	Pro 20	320	Asn	Ile	_'ys	Gln 25	عت:	Ile	Ala	Leu	Leu 30	Gln	Arg
15	Gly	Ast.	Cys 35	Thr	Phe	Lys	Glu	Lys 40	Ile	Ser	Arg	Ala	Ala 45	Phe	His	Asn
20	a <u>·</u> a	Val 50	Ala	Val	Val	Ile	Τ⁄τ 55	Asn	Asn	Lys	Sar	Lys 60	Glu	Glu	Pro	Val
-0	Thr 65	Met	T.L.	His	Pro	GLy TO	Thr	Glu	His	Ile	Ile 75		Val	Met	Ile	Thr 80
25	Glu	Leu	Arg	Gly	Lys 25	ζελ	Ile	Leu	Ser	77 <u>7</u> 90	Leu	Glu	Lys	Asn	Ile 95	
	Val	Glm	Mez	Thr 100	Ile	Α <u>`</u> a	Val	Зlу	Thr 105	Arg	Met	Pro	Pro	Lys 110	Asn	Phe
30	Ser	Yzâ	Gly 115	Ser	Læu	Val	Phe	Val 120	Ser	Ile	Ser	Phe	Ile 125	٧al	Leu	Met
35	lle	130	Ser	Ser	Ala	dzī	Leu 135	lle	?he	Tyr	Pine	Ile 140	Gln	Lys	Ile	Arg
	7yr 145	Thr	Asti	Ala	Arg	Asp 150	Arg	Asn	Gln	Arg	Arg 155	Leu	Gly	Asp	Ala	Ala 160
40	Lys	Lys	Ala	Ile	Ser 155	L∵s	Leu	Thr	Thr	Arg 170	Thr	Val	Lys	Lys	Gly 175	Asp
	Lys	Glu	The	Asp 180	P20	Asp	Phe	Asp	His 185	Cys	Ala	7al	Cys	Ile 190	Glu	Ser
45	<u>:72</u>	Lys	G <u>ln</u> 195	Asn	Asp	Val	Val	Arg 200	Ile	Leu	Pro	Cys	Lys 205	His	Val	Phe
50	His	Lуs 219	Ser	Cys	Val	Asp	Pro 215	grp	Гел	Ser	Glu	His 220	Cys	Thr	Cys	Pro
50	<u>Met</u> 225	Cys	Lys	Leu	Asn	Ile 230	Leu	Lys	Ala	ŗen	Gly 235	Ile	Val	Pro	Asn	Leu 240
55	Pro	Суз	The	qzA	Asn 245	Val	Ala	?he	Asp	Met 250	Glu	Arg	Leu	Thr	Arg 255	Thr
	Gln	Ala	Va <u> </u>	Asn 260	λ≃g	Arg	Ser	Ala	Leu 265	Gly	çex	Leu	Ala	Gly 270	Asp	Asn

Ser Leu Gly Leu Glu Pro Leu Arg Thr Ser Gly Ile Ser Pro Leu Pro

	275	280	285
5	Gln Asp Gly Glu Leu Thr Pro 290 295	Arg Thr Gly Glu Ile 300	Asn Ile Ala Val
	Thr Lys Glu Trp Phe Ile Ile 305 310	Ala Ser Phe Gly Leu 315	Leu Ser Ala Leu 320
10	Thr Leu Cys Tyr Met Ile Ile 325	Arg Ala Thr Ala Ser	Leu Asn Ala Asn 335
	Glu Val Glu Trp Phe 340	,	
15	·		
	(2) INFORMATION FOR SEQ ID	NO: 704:	
20	(B) TYPE: ami (D) TOPOLOGY:	0 amino acids no acid	<b>4:</b> .
25	His Gly Val Ala Asp His Leu 1 5	Gly Cys Asp Pro Gln	Thr Arg Phe Phe
30	Val Pro Pro Asn Ile Lys Gln 20	Trp Ile Ala Leu Leu 25	Gln Arg Gly Asn 30
	Cys Thr Phe Lys Glu Lys Ile	Ser Arg Ala Ala Phe 40	His Asn Ala Val 45
35	Ala Val Val Ile Tyr Asn Asn 50 55	Lys Ser Lys Glu Glu 60	
40	(2) INFORMATION FOR SEQ ID	NO: 705:	
	(B) TYPE: am:	14 amino acids no acid	
45	(D) TOPOLOGY (xi) SEQUENCE DESCRI	linear PTION: SEQ ID NO: 70	5:
	Met Ser Gly Gln Gly Leu Ala 1 5	Gly Phe Phe Ala Ser	Val Ala Met Ile 15
50	Cys Ala Ile Ala Ser Gly Ser 20	Glu Leu Ser Glu Ser 25	Ala Phe Gly Tyr 30
55	Phe Ile Thr Ala Cys Ala Val	Ile Ile Leu Thr Ile 40	Ile Cys Tyr Leu 45
	Gly Leu Pro Arg Leu Glu Phe 50 55		
60	Glu Gly Pro Gly Glu Gln Glu	Thr Lys Leu Asp Leu	Ile Ser Lys Gly

WO 98/54963 PCT/US98/11422

	Glu	Glu	Pro	Arg	Ala 85	Gly	Lys	Glu	Glu	Ser 90	Gly	Val	Ser	Val	Ser 95	
5	Ser	Gln	Pro	Thr 100	Asn	Glu	Ser	His	Ser 105	Ile	Lys	Ala	Ile	Leu 110	Lys	Asn
10	Ile	Ser	Val 115	Leu	Ala	Phe	Ser	Val 120	Cys	Phe	Ile	Phe	Thr 125	Ile	Thr	Ile
	Gly	Met 130	Phe	Pro	Ala	Val	Thr 135	Val	Glu	Val	Lys	Ser 140	Ser	Ile	Ala	Gly
15	Ser 145	Ser	Thr	Trp	Glu	Arg 150	Tyr	Phe	Ile	Pro	Val 155	Ser	Cys	Phe	Leu	Thr 160
	Phe	Asn	Ile	Phe	Asp 165	Trp	Leu	Gly	Arg	Ser 170	Leu	Thr	Ala	Val	Phe 175	Met
20	Trp	Pro	Gly	Lys 180	Asp	Ser	Arg	Trp	Leu 185	Pro	Ser	Trp	Xaa	Leu 190	Ala	Arg
25	Leu	Val	Phe 195	Val	Pro	Leu	Leu	Leu 200	Leu	Cys	Asn	Ile	Lys 205	Pro	Arg	Arg
	Tyr	Leu 210	Thr	Val	Val	Phe	Glu 215	His	Asp	Ala	Trp	Phe 220	Ile	Phe	Phe	Met
30	Ala 225	Ala	Phe	Ala	Phe	Ser 230	Asn	Gly	Tyr	Leu	Ala 235	Ser	Leu	Суз	Met	Cys 240
	Phe	Gly	Pro	Lys	Lys 245	Val	Lys	Pro	Ala	Glu 250	Ala	Glu	Thr	Ala	Glu 255	Pro
35	Ser	Trp	Pro	Ser 260	Ser	Cys	Val	Trp	Val 265	Trp	His	Trp	Gly	Leu 270	Phe	Ser
10	Pro	Ser	Cys 275	Ser	Gly	Gln	Leu	Cys 280	Asp	Lys	Gly	Trp	Thr 285	Glu	Gly	Leu
	Pro	Ala 290	Ser	Leu	Pro	Val	Cys 295	Leu	Leu	Pro	Leu	Pro 300	Ser	Ala	Arg	Gly
15	Asp 305	Pro	Glu	Trp	Ser	Gly 310	Gly	Phe	Phe	Phe						
-0	(2)	INFO	ORMAT	rion	FOR	SEQ	ID N	ю: 7	06:						•	
50		٠	(i) s					ERIST 06 ar			is					
55		,	(xi)	(1	) T	OPOL	OGY:	no ad line PTION	ear	EQ II	O NO:	706	i:			
	Met 1	Ser	Gly	Gln	Gly 5	Leu	Ala	Gly	Phe	Phe 10	Ala	Ser	Val	Ala	Met 15	Ile
50	Суз	Ala	Ile	Ala	Ser	Gly	Ser	Glu	Leu	Ser	Glu	Ser	Ala	Phe	Gly	Tyr

				20					25		٠			30		
5	Phe	Ile	Thr 35	Ala	Cys	Ala	Val	Ile 40	Ile	Leu	Thr	Ile	Ile 45	Cys	Tyr	Leu
ر	Gly	Leu 50		Arg	Leu	Glu	Phe 55	Tyr	Arg	Tyr	Tyr	Gln 60	Gln	Leu	Lys	Leu
10	Glu 65		Pro	Gly	Glu	Gln 70	Glu	Thr	Lys	Leu	Asp 75	Leu	Ile	Ser	Lys	Gly 80
	Glu	Glu	Pro	Arg	Ala 85	Gly	Lys	Glu	Glu	Ser 90	Gly	Val	Ser	Val	Ser 95	Asn
15	Ser	Gln	Pro	Thr 100	Asn	Glu	Ser	His	Ser 105	Ile						
20	(2)	INF	ORMA:	rion	FOR	SEQ	ID I	NO: 7	707 :							
			(i)					ERIS.								
25	•		(xi)	(	B) T D) T	YPE : OPOL	ami OGY:	1 am no a lin PTIO	cid ear			: 70	7 :			
20	Ser 1		Val	Ser	Val 5	Ser	Asn	Ser	Gln	Pro 10	Thr	Asn	Glu	Ser	His 15	Ser
30	Ile	Lys	Ala	Ile 20	Leu	Lys	Asn	Ile	Ser 25	Val	Leu	Ala	Phe	Ser 30	Val	Cys
35	Phe	Ile	Phe 35	Thr	Ile	Thr	Ile	Gly 40	Met	Phe	Pro	Ala	Val 45	Thr	Val	Glu
	Val	Lys 50	Ser	Ser	Ile	Ala	Gly 55	Ser	Ser	Thr	Trp	Glu 60	Arg	Tyr	Phe	Ile
40	Pro 65	Val	Ser	Cys	Phe	Leu 70	Thr	Phe	Asn	Ile	Phe 75	Asp	Trp	Leu	Gly	Arg 80
	Ser							٠								÷
45																
	(2)	INF	ORMAT	rion	FOR	SEQ	ID 1	vo: 7	08:							٠
50 .			(i) :	(:	A) L B) T	engt Ype:	H: 9 ami	2 am no a	ino a		s .	•				
55			(xi)		-			line PTION		EQ II	ONO:	: 708	3:			
<i>33</i>	Thr 1	Ile	Gly	Met	Phe 5	Pro	Ala	Val	Thr	Val 10	Glu	Val	Lys	Ser	Ser 15	Ile
60	Ala	Gly	Ser	Ser 20	Thr	Trp	Glu	Arg	Тут 25	Phe	Ile	Pro	Val	Ser 30	Cys	Phe

	Leu	Thr	Phe 35	Asn	Ile	Phe	Asp	Trp 40	Leu	Gly	Arg	Ser	Leu 45	Thr	Ala	Val
5	Phe	Met 50	Trp	Pro	Gly	Lys	Asp 55	Ser	Arg	Trp	Leu	Pro 60	Ser	Trp	Xaa	Leu
10	Ala 65	Arg	Leu	Val	Phe	Val 70	Pro	Leu	Leu	Leu	Leu 75	Cys	Asn	Ile	Lys	Pro 80
	Arg	Arg	Tyr	Leu	Thr 85	Val	Val	Phe	Glu	His 90	Asp	Ala				
15	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	vo: 7	709:						٠	
20			(i) :	(	A) L	ENGT	H: 7	ERIS 4 am no a	ino		s					
				(	D) T	OPOL	OGY:	lin	ear							
			(xi)	SEQ	UENC	E DE:	SCRI	PTIO	N: S	EQ II	D NO	: 70	€:			
25	Phe 1	Gly	Pro	Lys	Lys 5	Val	Lys	Pro	Ala	Glu 10	Ala	Glu	Thr	Ala	Glu 15	Pro
	Ser	Trp	Pro	Ser 20	Ser	Cys	Val	Trp	Val 25	Trp	His	Trp	Gly	Leu 30	Phe	Ser
30	Pro	Ser	Суs 35	Ser	Gly	Gln	Leu	Cys 40	Asp	Lys	Gly	Trp	Thr 45	Glu	Gly	Leu
35	Pro	Ala 50	Ser	Leu	Pro	Val	Cys 55	Leu	Leu	Pro	Leu	Pro 60	Ser	Ala	Arg	Gly
	Asp 65	Pro	Glu	Trp	Ser	Gly 70	Gly	Phe	Phe	Phe						
40																
	(2)	INF	ORMAT	NOI	FOR	SEQ	ID N	10: 7	710:							
			(i) :					ERIST								
45								35 a no a		acı	as		•			
			(xi)					line PTIO		EQ II	O NO	: 710	):			
	Aso	Asn	Asp	Glv	Phe	Glu	Tle	Val	Pro	Tle	ćlu	1en	Pro	Δla	Tare	Hic
50	1			02,	5					10	014				15	*****
	Arg	Ile	Leu	Asp 20	Pro	Glu	Gly	Leu	Ala 25	Leu	Gly	Ala	Val	Ile 30	Ala	Ser
55	Ser	Lys	Lys 35	Ala	Lys	Arg	Asp	Leu 40	Ile	Asp	Asn	Ser	Phe 45	Asn	Arg	Тут
60	Thr	Phe 50	Asn	Glu	Asp	Glu	Gly 55	Glu	Leu	Pro	Glu	Trp 60	Phe	Val	Gln	Glu

	Glu 65	Lys	Gln	His	Arg	Ile 70	Arg	Gln	Leu	Pro	Val 75	Gly	Lys	Lys	Glu	Val 80
5	Glu	His	Tyr	Arg	Lys 85	Arg	Trp	Arg.	Glu 	Ile 90	Aşn	Ala	Arg	Pro	Ile 95	Xaa
	Xaa	Xaa	Xaa	Xaa 100	Xaa	Xaa	Xaa	Xaa	Xaa 105	Xaa	Xaa	Xaa	Xaa	Xaa 110	Xaa	Xaa
10	Leu	Glu	Gln 115	Thr	Arg	Lys	Lys	Ala 120	Glu	Àla	Val	Val	Asn 125	Thr	Val	Asp
15	Ile	Хаа 130	Arg	Thr	Arg	Glu	Ser 135									
	(2)	INF	ORMA	rion	FOR	SEQ	ID 1	vo: 7	711:							
20			(i) :	(	ENCE A) L B) T D) T	ENGT YPE :	H: 5 ami	0 am	ino d		s					
25			(xi)							EQ II	ON C	: 71	l:			
	Asp 1	Asp	Asp	Gly	Phe 5	Glu	Ile	Val	Pro	Ile 10	Glu	Asp	Pro	Ala	Lys 15	His
30	Arg	Ile	Leu	Asp 20	Pro	Glu	Gly	Leu	Ala 25	Leu	Gly	Ala	Val	Ile 30	Ala	Ser
	Ser	Lys	Lys 35	Ala	Lys	Arg	Asp	Leu 40	Ile	Asp	Asn	Ser	Phe 45	Asn	Arg	Tyr
35	Thr	Phe 50								,						٠
40	(2)	INF	ORMA	rion	FOR	SEQ	ID 1	NO: 7	712:							
45			(i) :	(	ENCE A) L B) T	ENGT YPE :	H: 5 ami	1 am	ino a		s					
			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: SI	EQ I	OM C	: 71	2:			
50	Lys 1	Arg	Trp	Arg	Glu 5	Ile	Asn	Ala	Arg	Pro 10	Ile	Xaa	Xaa	Xaa	Хаа 15	Xaa
•	Xaa	Xaa	Xaa	Хаа 20	Xaa	Xaa	Xaa	Xaa	Xaa 25	Xaa	Xaa	Xaa	Leu	Glu 30	Gln	Thr
55	Arg	Lys	Lys 35	Ala	Glu	Ala	Val	Val 40	Asn	Thr	Val	Asp	Ile 45	Xaa	Arg	Thr
	Arg	Glu 50	Ser													

	(2)	INF	ORMA'	PION	FOR	SEQ	ID I	NO:	713:							
5		•		(	A) L B) T D) T	ENGT YPE : OPOL	H: 2 ami OGY:	16 a no a lin	mino cid ear	aci		: 71	3:			
10	Met 1	Ile		Asp										Ser	Ser 15	Gln
15	Pro	Ala	His	Leu 20	Cys	Pro	Glu	Asn	Pro 25	Leu	Leu	His	Leu	Lys 30	Ala	Ala
,	Val	Lys	Glu 35	Lys	Lys	Arg	Asn	Lys . 40	Lys	Lys	Lys	Thr	Ile 45	Gly	Ser	Pro
20	Lys	Arg 50	Ile	Gln	Ser	Pro	Leu 55	Asn	Asn	Lys	Leu	Leu 60	Asn	Ser	Pro	Ala
	Lys 65	Thr	Leu	Pro	Gly	Ala 70	Cys	Gly	Ser	Pro	Gln 75	Lys	Leu	Ile	Asp	Gly 80
25	Phe	Leu	Lys	His	Glu 85	Gly	Pro	Pro	Ala	Glu 90	Lys	Pro	Leu	Glu	Glu 95	Leu
30	Ser	Ala	Ser	Thr 100	Ser	Gly	Val	Pro	Gly 105	Leu	Ser	Ser	Leu	Gln 110	Ser	Asp
	Pro	Ala	Gly 115	Cys	Val	Arg	Pro	Pro 120	Ala	Pro	Asn	Leu	Ala 125	Gly	Ala	Val
35	Glu	Phe 130	Asn	Asp	Val	Lys	Thr 135	Leu	Leu	Arg	Glu	Trp 140	Ile	Thr	Thr	Ile
	Ser 145	Asp	Pro	Met.	Glu	Glu 150	Aşp	Ile	Leu	Gln	Val 155	Val	Lys	Tyr	Cys	Thr 160
40	Asp	Leu	Ile	Glu	Glu 165	Lys	Asp	Leu	Glu	Lys 170	Leu	Asp	Leu	Val	Ile 175	Lys
45	Tyr	Met	Lys	Arg 180	Leu	Met	Gln	Gln	Ser 185	Val	Glu	Ser	Val	Trp 190	Asn	Met
	Ala	Phe	Asp 195	Phe	Ile	Leu		Asn 200		Gln	Val	Val	Leu 205	Gln	Gln	Thr
50	Tyr	Gly 210	Ser	Thr	Leu	Lys	Val 215	Thr								
55	(2)			rion				•								
			14/	SEQUI .)		CHAI ENGT					s					
						YPE:										

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 714:

**60** .

	Met 1	Ile	Lys	Asp	Lys 5	Gly	Arg	Ala	Arg	Thr 10	Ala	Leu	Thr	Ser	Ser 15	Gln
5	Pro	Ala	His	Leu 20	Cys	Pro	Glu	Asn	Pro 25	Leu	Leu	His	Leu	Lys 30	Ala	Ala
10	Val	Lys	G1u 35	Lys	Lys	Arg	Asn	Lys 40	Lys	Lys	Lys	Thr	Ile 45	Gly	Ser	Pro
	Lys	Arg 50	Ile	Gln									٠			
15	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	<b>1</b> 0: 1	715:						٥	
20			(i) :	(	A) L B) T	CHAI ENGT YPE: OPOL	H: 1 ami	00 a no a	mino cid		ds					
			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	: 71	5 :			
25	Lys 1	Arg	Ile	Gln	Ser 5	Pro	Leu	Asn	Asn	Lys 10	Leu	Leu	Asn	Ser	Pro 15	Ala
	Lys	Thr	Leu	Pro 20	Gly	Ala	Cys	Gly	Ser 25	Pro	Gln	Lys	Leu	Ile 30	Asp	Gly
30	Phe	Leu	Lys 35	His	Glu	Gly	Pro	Pro 40	Ala	Glu	Lys	Pro	Leu 45	Glu	Glu	Leu
35	Ser	Ala 50	Ser	Thr	Ser	Gly	Val 55	Pro	Gly	Leu	Ser	Ser 60	Leu	Gln	Ser	Asp
	Pro 65	Ala	Gly	Cys	Val	Arg 70	Pro	Pro	Ala	Pro	Asn 75	Leu	Ala	Gly	Ala	Val 80
40	Glu	Phe	Asn	Asp	Val 85	Lys	Thr	Leu	Leu	Arg 90	Glu	Trp	Ile	Thr	Thr 95	Ile
_	Ser	Asp	Pro	Met 100												
45																
	(2)	INF	ORMA	TION	FOR	SEQ	ID I	NO:	716:					•		
50			(i)	(	A) I B) I	CHA ENGI YPE:	H: 7	4 an	nino cid		ls			•		
			(xi)	SEQ						EQ I	D NO	: 71	6:			
55	Thr 1		Ser	Asp	Pro 5		Glu	Glu	Asp	Ile 10	Leu	Gln	Val	Val	Lys 15	Tyr
60	Cys	Thr	Asp	Leu 20	Ile	Glu	Glu	Lys	Asp 25	Leu	Glu	Lys	Leu	Asp 30	Leu	Val

(2) INFORMATION FOR SEQ ID NO: 720:

```
Ile Lys Tyr Met Lys Arg Leu Met Gln Gln Ser Val Glu Ser Val Trp
      Asn Met Ala Phe Asp Phe Ile Leu Asp Asn Val Gln Val Val Leu Gln
 5
      Gln Thr Tyr Gly Ser Thr Leu Lys Val Thr
                          70
10
      (2) INFORMATION FOR SEQ ID NO: 717:
             (i) SEQUENCE CHARACTERISTICS:
15
                    (A) LENGTH: 18 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 717:
20
      Phe Cys His Asp Cys Lys Phe Pro Glu Ala Ser Pro Ala Met Asn Cys
      Glu Pro
25
      (2) INFORMATION FOR SEQ ID NO: 718:
30
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 18 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 718:
35
      Phe Cys His Asp Cys Lys Phe Pro Glu Ala Ser Pro Ala Met Asn Cys
                       5
                                           10
      Glu Pro
40
      (2) INFORMATION FOR SEQ ID NO: 719:
45
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 27 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
50
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 719:
      Pro Gln Pro Ser Asn Phe Pro Thr Thr Val Arg Asn Leu Pro Tyr Ser
                  5
                                          10
55
      Gly Ala Gly Ala Gln Pro Pro Pro Ser Asn Cys
                                     25
```

			(i) :			CHAI ENGT					ds					
5			(xi)	(	D) T	YPE: OPOL E DE:	OGY:	lin	ear	EQ II	D NO	: 72	0:			•
10	Met 1	Ala	Ser	Ser	Val 5	Pro	Ala	Gly	Gly	His 10	Thr	Arg	Ala	Gly	Gly 15	Ile
10	Phe	Leu	Ile	Gly 20	Lys	Leu	Asp	Leu	Glu 25	Ala	Ser	Leu	Phe	Lys 30	Ser	Phe
15	Gln	Trp	Leu 35	Pro	Phe	Val	Leu	Arg 40	Lys	Lys	Cys	Asn	Phe 45	Phe	Cys	Trp
	Asp	Ser 50	Ser	Ala	His	Ser	Leu 55	Pro	Leu	His	Pro	Leu 60	Ser	Ala	Ser	Cys
20	Ser 65	Ala	Pro	Ala	Cys	His 70	Ala	Ser	Asp	Thr	His 75		Leu	Tyr	Pro	Ser 80
25	Thr	Arg	Ala	Leu	Cys 85	Pro	Ser	Ile	Phe	Ala 90	Trp	Leu	Val	Ala	Pro 95	His
•	Ser	Val	Phe	Arg 100		Asn	Ala	Pro	Gly 105	Pro	Thr	Pro	Ser	Ser 110	Gln	Ser
30	Ser	Pro	Val 115		Pro	Val	Phe	Pro 120	Val	Ser	Phe	Met	Ala 125	Leu	Ile	Val
	Cys	Xaa 130	Leu	Val	Cys	Cys						•				
35	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: '	721:							
40				(	A) L B) T D) T	CHA ENGT YPE: OPOL E DE	H: 7 ami OGY:	1 am no a lin	ino cid ear	acid		: 72:	1:			
45	Met	Ala	Ser	Ser	Val 5	Pro	Ala	Gly	Gly	His 10	Thr	Arg	Ala	Gly	Gly 15	Ile
50	Phe	Leu	Ile	Gly 20	Lys	Leu	Asp	Leu	Glu 25	Ala	Ser	Leu	Phe	Lys 30	Ser	Phe
	Gln	Trp	Leu 35	Pro	Phe	Val	Leu	Arg 40	Lys	Lys	Cys	Asn	Phe 45	Phe	Cys	Trp
55	Asp	Ser 50	Ser	Ala	His	Ser	Leu 55	Pro	Leu	His	Pro	Leu 60	Ser	Ala	Ser	Cys
	Ser 65	Ala	Pro	Ala	Cys	His 70	Ala									

712

	(2)	INF	ORMA!	rion	FOR	SEQ	ID I	NO: 1	722:							
5				(	A) L B) T D) T	ENGT YPE: OPOL	H: 4 ami OGY:	6 am no a lin	ino cid ear	acid		: 72	2:			
10	Phe 1	Ala	Trp	Leu	Val 5	Ala	Pro	His	Ser	Val 10	Phe	Arg	Thr	Asn	Ala 15	Pro
15	Gly	Pro	Thr	Pro 20	Ser	Ser	Gln	Ser	Ser 25	Pro	Val	Phe	Pro	Val 30	Phe	Pro
	Val	Ser	Phe 35	Met	Ala	Leu	Ile	Val 40	Cys	Xaa	Leu	Val	Cys 45	Суѕ		
20	(2)	INFO	ORMA:	rion	FOR	SEQ	ID 1	NO: 7	723:						*	
25				(	A) L B) T D) T	ENGT YPE: OPOL	H: 1 ami OGY:	34 a no a lin	mino cid ear	aci		: 72.	3:			•
80	Met 1	Ala	Ser	Ser	Val 5	Pro	Ala	Gly	Gly	His 10	Thr	Arg	Ala	Gly	Gly 15	Ile
	Phe	Leu	Ile	Gly 20	Lys	Leu	Asp	Leu	Glu 25	Ala	Ser	Leu	Phe	Lys 30	Ser	Phe
35	Gln	Trp	Leu 35	Pro	Phe	Val	Leu	Arg 40	Lys	Lys	Cys	Asn	Phe 45	Phe	Cys	Trp
10	Ąsp	Ser 50	Ser	Ala	His	Ser	Leu 55	Pro	Leu	His	Pro	Leu 60	Ser	Ala	Ser	Cys
	Ser 65	Ala	Pro	Ala	Cys	His 70	Ala	Ser	Asp	Thr	His 75		Leu	Tyr	Pro	Ser 80
15	Thr	Arg	Ala	Leu	Cys 85	Pro	Ser	Ile	Phe	Ala 90	Trp	Leu	Val	Ala	Pro 95	His
	Ser	Val	Phe	Arg 100	Thr	Asn	Ala	Pro	Gly 105	Pro	Thr	Pro	Ser	Ser 110	Gln	Ser
50	Ser	Pro	Val 115	Phe	Pro	Val	Phe	Pro 120	Val	Ser	Phe	Met	Ala 125	Leu	Ile	Val
55	Cys	Xaa 130	Leu	Val	Cys	Cys		٠								
	(2)	TAICY	1DMA	PTON	FOR	CEO	TD 3		724.							

(i) SEQUENCE CHARACTERISTICS:

				. (	B) I D) I	YPE: OPOL	ami OGY:	no a lin	cid ear	aci						
5			(xi)						*	_						
	Met 1	Ala	Met	Glu	Gly 5	Tyr	Trp	Arg	Phe	Leu 10	Ala	Leu	Leu	Gly	Ser 15	Ala
10 -	Leu	Leu	Val	Gly 20	Phe	Leu	Ser	Val	Ile 25	Phe	Ala	Leu	Val	Trp 30	Val	Lei
	His	Tyr	Arg 35	Glu	Gly	Leu	Gly	Trp 40	Asp	Gly	Ser	Ala	Leu 45	Glu	Phe	Asr
15	Trp	His 50	Pro	Val	Leu	Met	Val 55		Gly	Phe	Val	Phe 60	Ile	Gln	Glý	Ile
20	Ala 65	Ile	Ile	Val	Tyr	Arg 70	Leu	Pro	Trp	Thr	Trp 75	Lys	Cys	Ser	Lys	Let 80
	Leu	Met	Lys	Ser	Ile 85	His	Ala	Gly	Leu	Asn 90	Ala	Val	Ala	Ala	Ile 95	Leu
25	Ala	Ile	Ile	Ser 100	Val	Val	Ala	Val	Phe 105	Glu	Asn	His	Asn	Val 110	Asn	Ası
	Ile	Ala	Asn 115	Met	Tyr	Ser	Leu	His 120	Ser	Trp	Val	Gly	Leu 125	Ile	Ala	Va]
30	Ile	Cys 130	Tyr	Leu	Leu	Gln	Leu 135	Leu	Ser	Gly	Phe	Ser 140	Val	Phe	Leu	Leu
35	Pro 145	Trp	Ala	Pro	Leu	Ser 150	Leu	Arg	Ala	Phe	Leu 155	Met	Pro	Ile	His	Val
	Tyr	Ser	Gly	Ile	Val 165		Phe	Gly	Thr	Val 170	Ile	Ala	Thr	Alá	Leu 175	Met
40	Gly	Leu	Thr	Glu 180	Lys	Leu	Ile	Phe	Ser 185	Leu	Arg	Asp	Pro	Ala 190	Tyr	Ser
	Thr	Phe	Pro 195	Pro	Glu	Gly	Val	Phe 200	Val	Asn	Thr	Leu	Gly 205	Leu	Leu	Ile
45	Leu	Val 210	Phe	Gly	Ala	Leu	Ile 215	Phe	Trp	Ile	Val	Thr 220	Arg	Pro	Gln	Trp
50	Lys 225	Arg	Pro	Lys	Glu	Pro 230	Asn	Ser	Thr	Ile	Leu 235	His	Pro	Asn	Gly	Gly 240
50	Thr	Glu	Gln	Gly	Ala 245	Arg	Gly	Ser	Met	Pro 250	Ala	Tyr	Ser	Gly	Asn 255	Asn
55	Met	Asp	Lys	Ser 260	Asp	Ser	Glu	Leu	Asn 265	Ser	Glu	Val	Ala	Ala 270	Arg	Lys
	Arg	Asn	Leu 275	Ala	Leu	Asp	Glu	Ala 280	Gly	Gln	Arg	Ser	Thr 285	Met		

	(2) INFORMATION FOR SEQ ID NO: 725:
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 43 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 725:</li> </ul>
10	Pro Gly Arg Ala Gly Pro Ser Pro Gly Leu Ser Leu Gln Leu Pro Ala 1 5 10 15
15	Glu Pro Gly His Pro Ala Gly Asn Leu Ala Pro Leu Thr Ser Arg Pro 20 25 30
	Gln Pro Leu Cys Arg Ile Pro Ala Val Pro Gly 35 40
20	(2) INFORMATION FOR SEQ ID NO: 726:
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 424 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 726:
30	Met Lys Leu Leu Gly Glu Cys Ser Ser Ser Ile Asp Ser Val Lys Arg 1 5 10 15
	Leu Glu His Lys Leu Lys Glu Glu Glu Glu Ser Leu Pro Gly Phe Val 20 25 30
35	Asn Leu His Ser Thr Glu Thr Gln Thr Ala Gly Val Ile Asp Arg Trp 35 40 45
40	Glu Leu Cln Ala Cln Ala Leu Ser Lys Glu Leu Arg Met Lys Gln 50 55 60
	Asn Leu Gln Lys Trp Gln Gln Phe Asn Ser Asp Leu Asn Ser Ile Trp 65 70 75 80
45	Ala Trp Leu Gly Asp Thr Glu Glu Glu Leu Glu Gln Leu Gln Arg Leu  85 90 95
50	Glu Leu Ser Thr Asp Ile Gln Thr Ile Glu Leu Gln Ile Lys Lys Leu 100 105 110
30	Lys Glu Leu Gln Lys Ala Val Asp His Arg Lys Ala Ile Ile Leu Ser 115 120 125
55	Ile Asn Leu Cys Ser Pro Glu Phe Thr Gln Ala Asp Ser Lys Glu Ser 130 135 140
	Arg Asp Leu Gln Asp Arg Leu Xaa Gln Met Asn Gly Arg Trp Asp Arg 145 150 155 160
60	Val Cys Ser Leu Leu Glu Glu Trp Arg Gly Leu Leu Gln Asp Ala Leu 165 170 175

	Met	Gln	Cys	Gln 180	Gly	Phe	His	Glu	Met 185	Ser	His	Gly	Leu	Leu 190	Leu	Met
5	Leu	Glu	Asn 195	Ile	Asp	Arg	Arg	Lys 200	Asn.	Glu	Ile	Val	Pro 205	Ile	Asp	Ser
10	Asn	Leu 210	Asp	Ala	Glu	Ile	Leu 215	Gln	Asp	His	His	Lys 220	Gln	Leu	Met	Gln
	Ile 225	Lys	His	Glu	Leu	Leu 230	Glu	Ser	Gln	Leu	Arg 235	Val	Ala	Ser	Leu	Gln 240
15	Asp	Met	Ser	Cys	Gln 245	Leu	Leu	Val	Asn	Ala 250	Glu	Gly	Thr	Asp	Суs 255	Leu
•	Glu	Ala	Lys	Glu 260	Lys	Val	His	Val	Ile 265	Gly	Asn	Arg	Leu	Lys 270	Leu	Leu
20	Leu	Lys	Glu 275	Val	Ser	Arg	His	Ile 280	Lys	Glu	Leu	Glu	Lys 285	Leu	Leu	Asp
25	Val	Ser 290	Ser	Ser	Gln	Gln	Asp 295	Leu	Ser	Ser	Trp	Ser 300	Ser	Ala	Asp	Glu
	Leu 305	Asp	Thr	Ser	Gly	Ser 310	Val	Ser	Pro	Xaa	Ser 315	Gly	Arg	Ser	Thr	Pro 320
30	Asn	Arg	Gln	Lys	Thr 325	Pro	Arg	Gly	Lys	Cys 330	Ser	Leu	Ser	Gln	Pro 335	Gly
	Pro	Ser	Val	Ser 340	Ser	Pro	His	Ser	Arg 345	Ser	Thr	Lys	Gly	Gly 350	Ser	Asp
35	Ser	Ser	Leu 355	Ser	Glu	Pro	Xaa	Pro 360	Gly	Arg	Ser	Gly	Arg 365	Gly	Phe	Leu
10	Phe	Arg 370	Val	Leu	Arg	Ala	Ala 375	Leu	Pro	Leu	Gln	Leu 380	Leu	Leu	Leu	Leu
	Leu 385	Ile	Gly	Leu	Ala	Суs 390	Leu	Val	Pro	Met	Ser 395	Glu	Glu	Ąsp	Тут	Ser 400
15	Cys	Ala	Leu	Ser	Asn 405	Asn	Phe	Ala	Arg	Ser 410	Phe	His	Pro	Met	415	Arg
	Tyr	Thr	Asn	Gly 420	Pro	Pro	Pro	Leu								
50																
	(2)	INFO	ORMAT	CION	FOR	SEQ	ID N	NO: 7	27:							
55			(i) :	(:	A) L B) T	engt: Ype:	H: 1	ERIST 10 am no ac line	mino cid		is .	•				
			(xi)	SEQU	JENCI	E DES	SCRI	PTION	V: SI	EQ II	) NO:	: 727	<b>7</b> :			
60	Met	Lys	Leu	Leu	Gly	Glu	Cys	Ser	Ser	Ser	Ile	Asp	Ser	Val	Lys	Arg

	1				5					10					15	
5	Leu	Glu	His	Lys 20	Leu	Lys	Glu	Glu	Glu 25	Glu	Ser	Leu	Pro	Gly 30	Phe	Val
	Asn	Leu	His 35	Ser	Thr	Glu	Thr	Gln 40	Thr	Ala	Gly	Val	Ile 45	Asp	Arg	Trp
10	Glu	Leu 50	Leu	Gln	Ala	Gln	Ala 55	Leu	Ser	Lys	Glu	Leu 60	Arg	Met	Lys	Gln
	Asn 65	Leu	Gln	Lys	Trp	Gln 70	Gln	Phe	Asn	Ser	Asp 75	Leu	Asn	Ser	Ile	Trp 80
15	Ala	Trp	Leu	Gly	Asp 85	Thr	Glu	Glu	Glu	Leu 90	Glu	Gln	Leu	Gln	Arg 95	Leu
20	Glu	Leu	Ser	Thr 100	Asp	Ile	Gln	Thr	Ile 105	Glu	Leu	Gln	Ile	Lys 110		٠
	(2)	INFO	ORMA!	rion	FOR	SEQ	ID 1	vo: 7	728:							
25			(i)	(	A) L B) T	ENGT YPE :	H: 1 ami	36 a no a	mino cid		đs					
30			(xi)	SEQ	D) T UENC					EQ I	оис	: 72	8:			
	Lys 1	Leu	Lys	Glu	Leu 5	Gln	Lys	Ala	Val	Asp 10	His	Arg	Lys	Ala	Ile 15	Ile
35	Leu	Ser	Ile	Asn 20	Leu	Cys	Ser	Pro	Glu 25	Phe	Thr	Gln	Ala	Asp 30	Ser	Lys
	Glu	Ser	Arg 35	Asp	Leu	Gln	Asp	Arg 40	Leu	Xaa	Gln	Met	Asn 45	Gly	Arg	Trp
40	Asp	Arg 50	Val	Суз	Ser	Leu	Leu 55	Glu	Glu	Trp	Arg	Gly 60	Leu	Leu	Gln	Asp
45	Ala 65	Leu	Met	Gln	Cys	Gln 70	Gly	Phe	His	Glu	Met 75	Ser	His	Gly	Leu	Leu 80
	Leu	Met	Leu	Glu	Asn 85	Ile	Asp	Arg	Arg	Lys 90	Asn	Glu	Ile	Val	Pro 95	Ile
50	Asp	Ser	Asn	Leu 100	Asp	Ala	Glu	Ile	Leu 105	Gln	Asp	His	His	Lys 110	Gln	Leu
	Met	Gln	Ile 115	Lys	His	Glu	Leu	Leu 120	Glu	Ser	Gln	Leu	Arg 125	Val	Ala	Ser
55	Leu	Gln 130	Asp	Met	Ser	Cys	Gln 135	Leu								÷
			-								• ,					

5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 105 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear															
			(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: 5	EQ I	D NO	: 72	9:			
10	Gln 1	Asp	Met	Ser	Cys 5	Gln	Leu	Leu	Val	Asn 10	Ala	Glu	Gly	Thr	Asp 15	Суз
•	Leu	Glu	Ala	Lys 20	Glu	Lys	Val	His	Val 25	Ile	Gly	Asn	Arg	Leu 30	Lys	Leu
15	Leu	Leu	Lys 35	Glu	Val	Ser	Arg	His 40	Ile	Lys	Glu	Leu	Glu 45	Lys	Leu	Leu
	Asp	Val 50	Ser	Ser	Ser	Gln	Gln 55	Asp	Leu	Ser	Ser	Trp 60	Ser	Ser	Ala	Asp
20	Glu 65	Leu	Asp	Thr	Ser	Gly 70	Ser	Val	Ser	Pro	Xaa 75	Ser	Gly	Arg	Ser	Thr 80
25	Pro	Asn	Arg	Gln	Lys 85	Thr	Pro	Arg	Gly	Lys 90	Cys	Ser	Leu	Ser	Gln 95	Pro
	Gly	Pro	Ser	Val 100	Ser	Ser	Pro	His	Ser 105						)	
30	(2)	INF	ORMA!	rion	FOR	SEQ	ID 1	NO: 7	730:				•	٠		
			(i)	-												
35				(	в) т	YPE:	ami	3 am no a lin	cid	acid	S					
			(xi)							EQ I	D NO	: 73	0:			
40	Asp 1	Ser	Ser	Leu	Ser 5	Glu	Pro	Xaa	Pro	Gly 10	Arg	Ser	Gly	Arg	Gly 15	Phe
	Leu	Phe	Arg	Val 20	Leu	Arg	Ala	Ala	Leu 25	Pro	Leu	Gln	Leu	Leu 30	Leu	Leu
45	Leu	Leu	Ile 35	Gly	Leu	Ala	Cys	Leu 40	Val	Pro	Met	Ser	Glu 45	Glu	Asp	Тут
50	Ser	Суs 50	Ala	Leu	Ser	Asn	Asn 55	Phe	Ala	Arg	Ser	Phe .60	His	Pro	Met	Leu
	Arg 65	Tyr	Thr	Asn	Gly	Pro 70	Pro	Pro	Leu							
55	(2)	INF	ORMA	rion	FOR	SEQ	ID 1	NO: 7	731:							
			(i):													
60								6 am no a		acid	s					

(B) TYPE: amino acid

•			(xi)					: lii		SEQ 1	D NO	): 7:	31:			
5	Met		. Leu	. Leu	Ile 5		Gly	/ Asn	Туг	Leu 10		Pro	Ser	His	Ser 15	Glu
	Ser	Ser	Arg	Arg 20		Cys	Leu	. Leu	Cys 25		Тух	Pro	Leu	Cys 30		Glu
10	Ile	Asn	Phe 35		Met	Lys	Val	Phe 40		Ser	Met	Pro	Phe 45		Val	Leu
15	Phe	Gln 50		Leu	Ile	Gln	Glu 55	Asp						·		
20	(2)	INF		SEQU	ENCE (A) I (B) I	CHA ENGI	RACT H: 2	NO: ERIS 271 a no a lin	TICS mind		.ds					
25	Arg 1							PTIC Ser						Lys	Tyr 15	Asp
30	Tyr	Leu	Pro	Thr 20	Thr	Val	Asn	Val	Cys 25	Ser	Glu	Leu	Val	Lys 30		Val
	Phe	Cys	Val 35	Leu	Val	Ser	Phe	Cys 40	Val	Ile	Lys	Lys	Asp 45	His	Gln	Ser
35	Arg	Asn 50	Leu	Lys	Tyr	Ala	Ser 55	Trp	Lys	Glu	Phe	Ser 60	Asp	Phe	Met	Lys
40	Trp 65	Ser	Ile	Pro	Ala	Phe 70	Leu	Tyr	Phe	Leu	Asp 75	Asn	Leu	Ile	Val	Phe 80
	Tyr	Val	Leu	Ser	Tyr 85	Leu	Gln	Pro	Ala	Met 90	Ala	Val	Ile	Phe	Ser 95	Asn
45	Phe	Ser	Ile	Ile 100	Thr	Thr	Ala	Leu	Leu 105	Phe	Arg	Ile	Val	Leu 110	Lys	Xaa
	Arg	Leu	Asn 115	Trp	Ile	Gln	Trp	Ala 120	Ser	Leu	Leu	Thr	Leu 125	Phe	Leu	Ser
50	Ile	Val 130	Ala	Leu	Thr	Ala	Gly 135	Thr	Lys	Thr	Leu	Gln 140	His	Asn	Leu	Ala
55	Gly 145	Arg	Gly	Phe	His	His 150	Asp	Ala	Phe	Phe	Ser 155	Pro	Ser	Asn	Ser	Cys 160
	Leu	Leu	Phe	Arg	Asn 165	Glu	Cys	Pro	Arg	Lys 170	Asp	Asn	Суз	Thr	Ala 175	Lys
	Glu	Trp	Thr	Phe	Pro	Glu	Ala	Lys	Trp	Asn	Thr	Thr	Ala	Arg	Val	Phe

	Sei	c His	195		Leu	Gly	Met	Gly 200		Val	Leu	Ile	Ile 205	Val	Gln	Cys
. 5	Phe	210	e Ser )	`Ser	Met		Asn 215	Ile	Tyr	Asn	Glu	Lys 220	Ile	Leu	Lys	Glu
10	Gl <sub>3</sub> 225		ı Gln	Leu	Thr	Glu 230	Xaa	Ile	Phe	Ile	Gln 235		Ser	Lys	Leu	Tyr 240
	Phe	e Phe	e Gly	Île	Leu 245	Phe	Asn	Gly	Leu	Thr 250	Leu	Gly	Leu	Gln	Arg 255	Ser
15	Asr	Arg	J Asp	Gln 260	Ile	Lys	Asn	Cys	Gly 265	Phe	Phe	Tyr	Gly	His 270	Ser	
20	(2)	INF	ORMA													
			(1)	(	ENCE A) L B) T	ENGT	н: 9	4 am	ino		s					,
25			(xi)		D) T					EQ I	D NO	: 73	3:			
	Asn 1		· Val	Pro	Asn 5	Leu	Gln	Thr	Leu	Ala 10	Val	Leu	Thr	Glu	Ala 15	Ile
30	Gly	Pro	Glu	Pro 20	Ala	Ile	Pro	Arg	Хаа 25	Pro	Arg	Glu	Pro	Pro 30	Val	Ala
35	Thr	Ser	Thr 35	Pro	Ala	Thr	Pro	Ser 40	Ala	Gly	Pro	Gln	Pro 45	Leu	Pro	Thr
	Gly	Thr 50	Val	Leu	Val	Pro	Gly 55	Gly	Pro	Ala	Pro	Pro 60	Cys	Leu	Gly	Glu
40	Ala 65		Ala	Leu	Leu	Leu 70	Pro	Pro	Cys	Arg	Pro 75	Ser	Leu	Thr	Ser	Cys 80
	Phe	Trp	Ser	Pro	Arg 85	Pro	Ser	Pro	Trp	Lys 90	Glu	Thr	Gly	Val		
45	. (2)	The	ODMA.	TON	<b>500</b>	~~~										
	(2)	INF	ORMAT													
50				(2 (I	A) LI 3) T	NGTI PE:	H: 40 amir	o amo	ino a		5					
`			(xi)		D) TO					Q II	NO:	734	:			
55	Ala 1	Leu	Gln	Leu	Ala 5	Phe	Tyr	Pro	Asp	Ala 10	Val	Glu (	Glu	Trp	Leu 15	Glu
60	Glu	Asn	Val	His 20	Pro	Ser	Leu	Gln	Arg 25	Leu	Gln :	Xaa :	Leu :	Leu 30	Gln .	Asp

WO 98/54963

720

Leu Ser Glu Val Ser Ala Pro Pro 35 5 (2) INFORMATION FOR SEQ ID NO: 735: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 30 amino acids 10 (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 735: Cys His Pro Pro Ala Leu Ala Gly Thr Leu Leu Arg Thr Pro Glu Gly 15 5 Arg Ala His Ala Arg Gly Leu Leu Glu Ala Gly Gly Ala 25 20 (2) INFORMATION FOR SEQ ID NO: 736: (i) SEQUENCE CHARACTERISTICS: 25 (A) LENGTH: 59 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 736: 30 Gly Ser Ser Ser Thr Arg Ser Trp Phe Ser Thr Ser Ser Pro Gln Arg Ser Ala Ser Trp His Ser Gly Ala Pro Ser Cys Arg Ser Trp Arg Leu 20 25 35 Pro Cys Ser Trp Leu Ser Thr Arg Met Pro Trp Arg Ser Gly Trp Arg 40 Lys Thr Cys Thr Pro Ala Cys Ser Gly Cys Lys 40 55 (2) INFORMATION FOR SEQ ID NO: 737: 45 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 247 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 737: Met Arg Pro Asp Trp Lys Ala Gly Ala Gly Pro Gly Gly Pro Pro Gln 55 Lys Pro Ala Pro Ser Ser Gln Arg Lys Pro Pro Ala Arg Pro Ser Ala 20 Ala Ala Ala Ile Ala Val Ala Ala Ala Glu Glu Glu Arg Arg Leu

•	Arg	Gln 50	. Arg	Asn	Arg	Leu	Arg 55		Glu	Glu	Asp	Lys 60		Ala	ı Val	. Glu
5	Arg 65	Cys	Leu	Glu	Glu	Leu 70	Val	Phe	Gly	' Asp	Val 75	Glu	Asn	Asp	Glu	Asp 80
	Ala	Leu	Leu	Arg	Arg 85	Leu	Arg	Gly	Pro	Arg 90	Val	Gln	Glu	His	Glu 95	Asp
10	Ser	Gly	Asp	Ser 100	Glu	Val	Glu	Asn	Glu 105		Lys	Gly	Asn	Phe 110		Pro
15	Gln	Lys	Lys 115	Pro	Val	Trp	Val	Asp 120	Glu	Glu	Asp	Glu	Asp 125	Glu	Glu	Met
	Val	Asp 130	Met	Met	Asn	Asn	Arg 135	Phe	Arg	Lys	Asp	Met 140	Met	Lys	Asn	Ala
20	Ser 145	Glu	Ser	Lys	Leu	Ser 150	Lys	Asp	Asn	Leu	Lys 155	Lys	Arg	Leu	Lys	Glu 160
	Glu	Phe	Gln	His	Ala 165	Met	Gly	Gly	Val	Pro 170	Ala	Trp	Ala	Glu	Thr 175	Thr
25	Lys	Arg	Lys	Thr 180	Ser	Ser	Asp	Asp	Glu 185	Ser	Glu	Glu	Asp	Glu 190	Asp	Asp
30	Leu	Leu	Gln 195	Arg	Thr	Gly	Asn	Phe 200	Ile	Ser	Thr	Ser	Thr 205	Ser	Leu	Pro
	Arg	Gly 210	Ile	Leu	Lys	Met	Lys 215	Asn	Cys	Gln	His	Ala 220	Asn	Ala	Glu	Arg
35	Pro 225	Thr	Val	Ala'	Arg	Ile 230	Ser	Ile	Cys	Ala	<b>Val</b> 235	Pro	Ser	Arg	Cys	Thr 240
	Asp	Cys	Asp		Cys 245	Trp	Asp	•								
40	(2)	INFO	RMAT	TON '	FOR	SEO	ID N	m . 7	30.				ē			
45	. •-•	(	i) S	EQUE A) B) C)	NCE  ) IN  ) TO	CHAF ENGTE (PE:	RACTE H: 18 amir XGY:	RIST 30 an 10 ac line	ICS: mino :id :ar	acio		. 738	:			
50	Cys 1	Leu (	Glu (	Glu 1	Leu 5	Val	Phe	Gly .	Asp '	Val (	Glu .	Asn i	Asp	Glu	Asp 15	Ala
55	Leu 1	Leu .	Arg i	Arg 1	Leu .	Arg	Gly	Pro l	Argʻ 25	Val (	Gln (	Glu 1	His	Glu 30	Asp	Ser
55	Gly 1	Asp :	Ser (	Glu V	/al (	Glu .	Asn (	Glu /	Ala i	Lys (	Gly A	Asn 1	Phe:	Pro	Pro (	Gln
60	Lys I	Lys 1 50	ero V	/al 1	, dz,	Val i	Asp (	Glu (	Glu <i>i</i>	Asp (	Glu A	Asp (	3lu (	Glu I	Met '	Val

	As 6	p Me 5	t Me	t Ası	n Ası	n Arg		e Arg	J Lys	s Asp	Met 75		t Lys	s Asr	ı Ala	a Ser 80
5	Gl	u Se	r Ly	s Lei	Sei 8		s Asp	) Ası	ı Let	1 Lys 90		s Ar	g Lei	ı Lys	95 95	
10	Ph	e Gl	n Hi	s Ala 100	a Met	t Gly	/ Gly	/ Val	105		Tr	Ala	a Glu	Thr		: Lys
	Arg	g Ly	s Thi	r Ser 5	: Sei	r Asp	Asp	120		r Glu	ı Glu	ı Ası	Glu 125		Asp	Leu
15	Le	130	n Ar	g Thr	: Gly	/ Asr	Phe 135		Ser	Thr	: Ser	Th:		Leu	Pro	Arg
	Gl <sub>3</sub> 145	7 Ile 5	e Le	ı Lys	Met	Lys 150		Cys	Gln	His	155		n Ala	Glu	Arg	Pro 160
20	Thi	Va.	l Ala	a Arg	Ile 165	e Ser	Ile	: Cys	Ala	Val 170		Ser	Arg	Суз	Thr 175	
25	Cys	s Ası	o Gly	/ Cys 180							,					
٠	(2)	INE	ORMA	TION	FOR	SEQ	ID:	NO:	∖ 739:							
30			(i)		(A) I (B) I	CHA LENGI TYPE:	H: 2	218 a no a	mino cid		.ds		-		,	
35	$\neg$			SEQ	UENC	E DE	SCRI	PTIO	N: S							
	1			Lys	5					10					15	
40	Phe	Leu	. Leu	Ile 20	Asn	Gly	Ile	Ala	Gly 25	Tyr	Leu	His	Leu	Leu 30	Ala	Met
	Lys	Thr	Lys 35	Glu	Leu	Ile	Gly	Ser 40	Met	Lys	Ile	Asn	Gly 45	Arg	Val	Ala
45	Ala	Ser 50	Thr	Phe	Ser	Ser	Asp 55	Ser	Lys	Lys	Val	<b>Tyr</b> 60	Ala	Ser	Ser	Gly
50	Asp 65	Gly	Glu	Val	Tyr	Val 70	Trp	Asp	Val	Asn	Ser 75	Arg	Lys	Cys	Leu	Asn 80
	Arg	Phe	Val	Asp	Glu 85	Gly	Ser	Leu	Tyr	Gly 90	Leu	Ser	Ile	Ala	Thr 95	Ser
55	Arg	Asn	Gly	Gln 100	Tyr	Val	Ala	Cys	Gly 105	Ser	Asn	Cys	Gly	Val 110	Val	Asn
	Ile	Tyr	Asn 115	Gln	Asp	Ser	Cys	Leu 120	Gln	Glu	Thr	Asn	Pro 125	Lys	Pro	Ile

	13	0			13	5		٠		14	0			•
5	Pro Th 145	r Thr	Glu	Ile L	eu Al 50	a Il	e Ala	a Sei	Glu 155		s Me	t Ly	s Gl	u Al 16
	Val Ar	g Leu	Val	His L 165	eu Pr	o Se	r Cys	Th:		. Phe	e Se	r As	n Ph 17	
10	Val II	e Lys	Asn 180	Lys A	sn Il	e Se:	r His		. His	Thi	. Me	t As <sub>l</sub>		e Se
•	Pro Ar	g Ser 195	Gly	Tyr P	he Ala	a Lei 200	u Gly O	/ Asn	Glu	Lys	Gly 209		s Ala	a Le
15	Met Tyr 210		Leu :	His H	is Tyr 21:		Asp	Phe						
20	(2) IN	FORMA	rion :	FOR SI	EQ ID	NO:	740:							
•		(i)			HARAC'				_			÷		
- 25		(xi)	(B (D	) TYP	GTH: E: am OLOGY DESCRI	ino a : lir	acid near			: 74	0 :	,		
30	Lys Ile	Asn	Gly ?	Arg Va	al Ala	Ala	Ser	Thr · 10	Phe	Ser	Ser	Asp	Ser 15	
	Lys Val	Tyr	Ala s 20	Ser Se	er Gly	Asp	Gly 25	Glu	Val	Tyr	Val	Trp 30		Val
35	Asn Ser	Arg 35	Lys (	∵ys Le	u Asn	Arg 40		Val	Asp	Glu	Gly 45		Leu	Tyr
	Gly Leu 50	Ser	Ile A	la Th	r Sęr 55	Arg	Asn	Gly	Gln	Тут 60	Val	Ala	Cys	Gly
40	Ser Asn 65	Cys	Gly V	al Va 7	l Asn O	Ile	Tyr	Asn	Gln 75	Asp	Ser	Cys	Leu	Gln 80
45	Glu Thr	Asn	Pro L	ys Pr 85	o Ile	Lys	Ala	Ile <sub>.</sub> 90	Met	Asn	Leu	Val	Thr 95	Gly
	Val Thr		100				105					110		
50	Ser Glu	Lys 1 115	Met L	ys Gl	u Ala	Val 120	Arg	Leu	Val	His	Leu 125	Pro	Ser	Cys
	Thr Val	Phe :	Ser A	sn Pho	Pro 135	Val	Ile	Lys		Lys 140	Asn	Ile	Ser	His
55	Val His 145	Thr 1	Met A	sp Phe 150	e Ser	Pro	Arg		Gly ' 155	Tyr	Phe	Ala	Leu	Gly 160
60	Asn Glu	Lys (		ys Ala 55 <sub>.</sub>	Leu							٠		

	12	) IN	FORM	ATTO	N FO	R SE	5 ID	NO:	741	:						
5	·			J	(A) (B) (D)	E CH LENG TYPE TOPO CE DI	TH: : am LOGY	246 ino : li:	amin acid near	o ac			4.1			
10	Met	: Ar	g Ile		ı Glı						a Leu			: Gly	/ Le	
15	Gly	/ G1:	y Glu	2 Thr 20	Arg	j Il∈	: Ile	Lys	Gly 25		e Glu	ı Cys	. Lys	Leu 30		s Sei
	Glr	Pr	o Trp 35	Glr	Ala	a Ala	Leu	Phe 40		Lys	Thr	Arg	Leu 45		ı Cys	Gly
20	Ala	Thu 50	r Leu )	l Ile	Ala	Pro	Arg 55	Trp	Leu	Leu	Thr	Ala 60		His	Cys	Let
25	Lys 65	Pro	Arg	Туг	Ile	Val 70	His	Leu	Gly	Gln	His 75		Leu	Gln	Lys	Glu 80
	Glu	Gly	/ Cys	Glu	Gln 85	Thr	Arg	Thr	Ala	Thr 90		Ser	Phe	Pro	His	
30	Gly	Ph€	e Asn	Asn 100	Ser	Leu	Pro	Asn	Lys 105	Asp	His	Arg	Asn	Asp 110		Met
	Leu	Val	. Lys 115	Met	Ala	Ser	Pro	Val 120	Ser	Ile	Thr	Trp	Ala 125	Val	Arg	Pro
35	Leu	Thr 130	Leu	Ser	Ser	Arg	Cys 135	Vál	Thr	Ala	Gly	Thr 140	Ser	Cys	Ser	Phe
40	Pro 145	Ala	Gly	Ala	Ala	Arg 150	Pro	Asp	Pro	Ser	Tyr 155	Ala	Cys	Leu	Thr	Pro 160
	Cys	Asp	Ala	Pro	Thr 165	Ser	Pro	Ser	Leu	Ser 170	Thr	Arg	Ser	Val	Arg 175	Thr
45	Pro	Thr	Pro	Ala 180	Thr	Ser	Gln	Thr	Pro 185	Trp	Cys	Val	Pro	Ala 190	Cys	Arg
	Lys	Gly	Ala 195	Arg	Thr	Pro	Ala	Arg 200	Val	Thr	Pro	Gly	Ala 205	Leu	Trp	Ser
50	Val	Thr 210	Ser	Leu	Phe	Lys	Ala 215	Leu	Ser	Pro	Gly	Ala 220	Arg	Ile	Arg	Val
55	Arg 225	Ser	Pro	Glu	Ser	Leu 230	Val	Ser	Thr	Arg	Lys 235	Ser	Ala	Asn	Met	Trp 240
	Thr	Gly	Ser	Arg	Arg 245	Arg										

	. (2	) IN	FORM	MIOITA	FOF	SEÇ	) ID	NO:	742	•						
5					(A) 1 (B) 1 (D) 1	LENG IYPE IOPO	TH: 2 : am: LOGY	228 a ino a : lir	amin acid near	S: o ac: SEQ ]		): <b>7</b> 4	12 :			
10	Glu 1	ı Thi	Arg	, Ile	Ile 5		Gly	Phe	Glu	Cys 10		Leu	His	Ser	Gln 15	
	Trp	Glr	Ala	Ala 20	Leu	Phe	Glu	Lys	Thr 25	Arg	Leu	Leu	Cys	Gly 30		Th
15	Leu	ı Ile	Ala 35	Pro	Arg	Trp	Leu	Leu 40	Thr	Ala	Ala	His	Cys 45		Lys	Pro
20	Arg	Tyr 50	Ile	Val	His	Leu	Gly 55	Gln	His	Asn	Leu	Gln 60		Glu	Glu	Gly
	Cys 65	Glu	Gln	Thr	Arg	Thr 70	Ala	Thr	Glu	Ser	Phe 75	Pro	His	Pro	Gly	Phe 80
25	Asn	. Asn	. Ser	Leu	Pro 85	Asn	Lys	Asp	His	Arg 90	Asn	Asp	Ile	Met	Leu 95	Va]
	Lys	Met	Ala	Ser 100	Pro	Val	Ser	Ile	Thr 105	Trp	Ala	Val	Arg	Pro 110	Leu	Thr
30	Leu	Ser	Ser 115	Arg	Cys	Val	Thr	Ala 120	Gly	Thr	Ser	Суз	Ser 125	Phe	Pro	Ala
35 ,	Gly	Ala 130	Ala	Arg	Pro	Asp	Pro 135	Ser	Tyr	Ala	Cys	Leu 140	Thr	Pro	Cys	Asp
	Ala 145	Pro	Thr	Ser	Pro	Ser 150	Leu	Ser	Thr	Arg	Ser 155	Val	Arg	Thr	Pro	Thr 160
40	Pro	Ala	Thr	Ser	Gln 165	Thr	Pro	Trp	Cys	Val 170	Pro	Ala	Cys	Arg	Lys 175	Gly
	Ala	Arg	Thr	Pro 180	Ala	Arg	Val	Thr	Pro 185	Gly	Ala	Leu	Trp	Ser 190	Val	Thr
45	Ser	Leu	Phe 195	Lys	Ala	Leu	Ser	Pro 200	Gly	Ala	Arg	Ile	Arg 205	Val	Arg	Ser
50	Pro	Glu 210	Ser	Leu	Val	Ser	Thr 215	Arg	Lys	Ser		Asn 220	Met	Trp	Thr	Gly
	Ser 225	Arg	Arg	Arg												

- (2) INFORMATION FOR SEQ ID NO: 743:
  - (i) SEQUENCE CHARACTERISTICS:
    - (A) LENGTH: 74 amino acids
- 60 (B) TYPE: amino acid

			(xi)		(D) I					EQ I	D NC	): 74	13:			
5	Cys 1	Lys	Leu	His	Ser 5	Gln	Pro	Trp	Gln	Aļa 10	Ala	Leu	Phe	Glu	Lys 15	Thr
	Arg	Leu	Leu	Cys 20	Gly	Ala	Thr	Leu	Ile 25	Ala	Pro	Arg	Trp	Leu 30	Leu	Thr
10	Ala	Ala	His 35	Cys	Leu	Lys	Pro	Arg 40	Tyr	Ile	Val	His	Leu 45	Gly	Glm	His
15	Asn	Leu 50	Gln	Lys	Glu	Glu	Gly 55	Cys	Glu	Gln	Thr	Arg 60	Thr	Ala	Thr	Glu
	Ser 65	Phe	Pro	His	Pro	Gly 70	Phe	Asn	Asn	Ser						
20	(2)	TNF	) RMA	ואסדיו	FOR	SEO	י דח	vo	744.							
25	(2)	· ·	(i) :	SEQUI ) ) )	ENCE A) L B) T D) T UENCE	CHA ENGT YPE: OPOL	RACT H: 8 ami OGY:	ERIS 1 am no a lin	TICS ino cid ear	acid		: 74	<b>4</b> :			
30	Val 1	Leu	Gln	Gly	Arg 5	Tyr	Phe	Ser	Pro	Ile 10	Leu	Glu	Met	Arg	Arg 15	Leu
	Arg	Pro	Glu	Gly 20	Xaa	Xaa	Asn	Leu	Pro 25	Gly	Gly	Ser	Arg	Ala 30	Gln	Lys
35	Glu	Pro	Arg 35	Gln	Asp	Leu	Thr	Leu 40	Val	Leu	Trp	Pro	His 45	C/s	Pro	His
40	Phe	Ala 50	Met	Thr	Arg	Ser	Tyr 55	Val	Pro	Thr	Lys	Gln 60	C'ys	Met	Val	31n
	Gly 65	Ser	Phe	Tyr	Cys	Ile 70	Phe	Ile	Phe	Lys	Gly 75	Pro	Val	Gln	Asn	80 GZĮ
45	Cys							•								
50	(2)			EQUE	FOR INCE	CHAF NGT!	ACTE	RIST	ICS:		ls					
55		Ċ	xi)	(1	B) TY D) TC ENCE	POL	GY:	line	ear	Q ID	NO:	745	i :			
	Met 1	Pro	Ile	Ile .	Asp (	3ln	Val .	Asn	Pro (	Glu 1 10	Leu :	His	Asp	Phe 1	Met (	Gln
60	Ser 2	Ala (	Glu '	Val (	Gly '	Thr	Ile	Phe .	Ala :	Leu :	Ser'	Trp	Leu	Ile '	Thr :	فترا

			20					25	5				30	)	
5	Phe Gl	y His 35	7a_	Leu	Ser	· Asp	Phe 40	e Arg	, His	7al	. Val	Arg 45		тут	Asp
	Phe Ph 5	e le: C	Ala	Cys	His	P#0 55	Leu	. Xet	Pro	Ile	T/r 50		Ala	Ala	Val
10	Ile Vai	l Lei	Tyr	Arg	Glu 70	Gln	`Glu	. Val	Leu	Asp 75		Asp	Cys	Asp	Met 80
	Ala Se	•		35	•				90	·				95	
15	Glu Thi		100					105					110		
20	Pro Asr	115					120					125			
	Glm Pro 130					135					140				_
25	Leu Ile 145				150					155					160
30	Asp Arg			155					170					175	
50	Lys Leu		120					135					190		
35	Ala Val Deu Phe 213	795	Lys	Ser	Ala	Leu	Glu 200	grb	Ala	Pro		Phe 205	Gln	Leu	Gln
40	(2) 그룹	ORMAT (i) s													
45		(xi)	(A (B (D	) LE ) TY ) TO	ingth Fe: Polo	: 70 amin GY:	ami o ac line	no a id ar	cids		746	:			
50	Cys Pro			5					10					15	
	Ala Phe		20					25					30		
55	Pro Gly	Gly I -35	eu A	la C	Sln A	Asn I	Leu 1 40	det 1	Pro I	Leu I	ors?	/al ( 45	Sly I	Phe :	ſrp
60	Met Gly 50	Ser L	eu P	ב סבי	3 oz	250 T	da,	ys 1	t dz.	Arg I	ys 1 60	rp V	al S	Ger (	Slų

```
Ala Cys Ser Cys Phe Cys
5
      (2) INFORMATION FOR SEQ ID NO: 747:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 20 amino acids
10
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 747:
     Gly Phe Gly Ser Val Ser Ala Ala Gly Arg Arg Ser Gly Gly Thr Trp
15
                                           10
      Gln Pro Val Gln
                   20
20
      (2) INFORMATION FOR SEQ ID NO: 748:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 16 amino acids
25
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 748:
      Pro Gly Gly Leu Ala Val Gly Ser Arg Trp Trp Ser Arg Ser Leu Thr
30
                                           10
        1
35
      (2) INFORMATION FOR SEQ ID NO: 749:
40
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 30 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 749:
45
      Leu Glu Pro Ser Arg Gln Arg Arg Pro Arg Arg Arg Gly Gly Thr Ser
      Arg Pro Glu Thr Asp Gln Arg Ala Lys Cys Trp Arg Gln Leu
50
                                        25
       (2) INFORMATION FOR SEQ ID NO: 750:
55
              (i) SEQUENCE CHARACTERISTICS:
                      (A) LENGTH: 11 amino acids
                      (B) TYPE: amino acid
                      (D) TOPOLOGY: linear
 60
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 750:
```

	Val (	Cys	Leu .	Arg	Cys ( 5	Gln .	Asn .	Arg 1	Met	Glu 10	Asn					
5																
	(2)	INFO	RMAT	ION	FOR	SEQ	ID N	0: 7	51:				•			
10			•	() () ()	INCE A) LE B) TV D) TO JENCE	INGTI (PE : OPOL(	i: 36 amir OGY:	57 an no ac line	mino cid ear	acio		: 751	.:			
15	Met .	Ala	Ala	Cys	Thr 5	Ala	Arg	Arg	Pro	Gly 10	Arg	Gly	Gln	Pro	Leu 15	Val
20	Val	Pro	Val	Ala 20	Asp	Xaa	Gly	Pro	Val 25	Ala	Lys	Ala	Ala	Leu 30	Cys	Ala
20	Ala	Хаа	Ala 35	Gly	Ala	Phe	Ser	Pro 40	Ala	Ser	Thr	Thr	Thr 45	Thr	Arg	Arg
25	His	Leu 50	Ser	Ser	Arg	Asn	Arg 55	Pro	Glu	Gly	Lys	Val 60	Leu	Glu	Thr	Val
	Gly 65	Val	Phe	Glu	Val	Pro 70	Lys	Gln	Asn	Gly	Lys 75	Tyr	Glu	Thr	Gly	Gln 80
30	Leu	Phe	Leu	His	Ser 85	Ile	Phe	Gly	Тут	Arg 90	Gly	Val	Val	Leu	Phe 95	Pro
35	Trp	Gln	Ala	Arg 100	Leu	Xaa	Asp	Arg	Asp 105	Val	Ala	Ser	Ala	Ala 110	Pro	Glu
	Lys	Ala	Glu 115	Asn	Pro	Ala		His 120	Gly	Ser	Lys	Glu	Val 125	Lys	Gly	Lys
40	Thr	His 130	Thr	Tyr	Tyr	Gln	Val 135		Ile	Asp	Ala	Arg 140	Asp	Cys	Pro	His
	Ile 145	Ser	Gln	Arg	Ser	Gln 150	Thr	Glu	Ala	Val	Thr 155		Leu	Ala	Asn	His 160
45	Asp	Asp	Ser	Arg	Ala 165		Tyr	Ala	Ile	Pro 170		Leu	Asp	Tyr	Val 175	Ser
50	His	Glu	. Asp	Ile 180		Pro	Tyr	Thr	Ser 185		Asp	Gln	Val	Pro 190	Ile	Gln
	His	Glu	Leu 195		Glu	Arg	Phe	Leu 200		Тут	Asp	Gln	Thr 205		Ala	Pro
55	Pro	Phe 210		Ala	Arg	Glu	Thr 215		Arg	Ala	Trp	Gln 220		Lys	Asn	His
	Pro 225		Leu	Glu	. Leu	Ser 230		Val	His	Arg	235		Thr	Glu	. Asn	1le 240
60	Arg	Val	Thr	Va]	. Ile	Pro	Phe	. Tyr	Met	: Gly	Met	. Arg	Glu	ı Ala	Gln	Asn

WO 98/54963 PCT/US98/11422

					245					250					255	
_	Ser	His	Val	Туг 260	Trp	Trp	Arg	Tyr	Cys 265	Ile	Arg	Leu	Glu	Asn 270	Leu	Asp
5	Ser	Asp	Val 275	Val	Gln	Leu	Arg	Glu 280	Arg	His	Trp	Arg	Ile 285	Phe	Ser	Leu
10	Ser	Gly 290	Thr	Leu	Glu	Thr	Val 295	Arg	Gly	Arg	Gly	Val 300	Val	Gly	Arg	Glu
	Pro 305		Leu	Ser	Lys	Glu 310	Gln	Pro	Ala	Phe	Gln 315	Tyr	Ser	Ser	His	Val 320
15	Ser	Leu	Gln	Ala	Ser 325	Ser	Gly	His	Met	Trp 330	Gly	Thr	Phe	Arg	Phe 335	Glu
20	Arg	Pro	Asp	Gly 340	Ser	His	Phe	Asp	Val 345	Arg	Ile	Pro	Pro	Phe 350	Ser	Leu
.20	Glu	Ser	Asn 355	Lys	Asp	Glu	Lys	Thr 360	Pro	Pro	Ser	Gly	Leu 365	His	Trp	
25	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	752:							
30					(A) I (B) T (D) T	ENGT TYPE :	TH: 3 ami	33 an ino a : lir	near	ació		): 75	·2:			
35	Met 1		Ala	Cys	Thr 5		Arg	Arg	Pro	Gly 10		Gly	Gln	Pro	Leu 15	Val
	Val	Pro	Val	. Ala		хаа	Gly	Pro	Val 25		Lys	Ala	Ala	Leu 30		Ala
40	Ala	٠.										•				
45	(2)	IN	FORMA	ATIOI	1 FOF	R SEÇ	) ID	NO:	753 :						•	
50				-	(A) : (B) : (D) :	LENG TYPE TOPO	TH: : am LOGY	33 a ino : li	STIC: mino acid near ON:	aci		0: <b>7</b> :	53:			
55	:	l			, !	5				10	0				19	
	Va.	l Pr	o Vai	l Ala 2		o Xaa	a Gl	y Pr	o Va. 2:		a Ly:	s Ala	a Ala	a Let 30		s Ala
60	Ala	a														

5	(2) INFORMATION FOR SEQ ID NO: 754:
3	<ul><li>(i) SEQUENCE CHARACTERISTICS:</li><li>(A) LENGTH: 33 amino acids</li><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li></ul>
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 754:
	Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val 1 5 10 15
15	Val Pro Val Ala Asp Xaa Gly Pro Val Ala Lys Ala Ala Leu Cys Ala 20 25 30
	Ala
20	
	(2) INFORMATION FOR SEQ ID NO: 755:
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 33 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
30	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 755:
	Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val  1 5 10 15
35	Val Pro Val Ala Asp Xaa Gly Pro Val Ala Lys Ala Ala Leu Cys Ala 20 25 30
	Ala
40	
	(2) INFORMATION FOR SEQ ID NO: 756:
45	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 33 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 756:</li> </ul>
50	Met Ala Ala Cys Thr Ala Arg Arg Pro Gly Arg Gly Gln Pro Leu Val
-	1 5 10 15
55	Val Pro Val Ala Asp Xaa Gly Pro Val Ala Lys Ala Ala Leu Cys Ala 20 25 30
	Ala

•	(2) INFORMATION FOR SEQ ID NO: 757:
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 35 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 757:</li> </ul>
10	Val Leu Glu Thr Val Gly Val Phe Glu Val Pro Lys Gln Asn Gly Lys  1 5 10 15
	Tyr Glu Thr Gly Gln Leu Phe Leu His Ser Ile Phe Gly Tyr Arg Gly 20 25 30
15	Val Val Leu 35
20	(2) INFORMATION FOR SEQ ID NO: 758:
25	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 16 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 758:</li> </ul>
30	Gly Leu Asp Tyr Val Ser His Glu Asp Ile Leu Pro Tyr Thr Ser Thr 1 5 10 15
35	(2) INFORMATION FOR SEQ ID NO: 759:
40	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 19 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 759:</li> </ul>
45	Asp Val His Arg Glu Thr Thr Glu Asn Ile Arg Val Thr Val Ile Pro 1 5 10 15
	Phe Tyr Met
50	
	(2) INFORMATION FOR SEQ ID NO: 760:  (i) SEQUENCE CHARACTERISTICS:
55	(A) LENGTH: 21 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 760:
60	Tro Tro Arg Tvr Cvs Ile Arg Leu Glu Asn Leu Asp Ser Asp Val Val

```
10
      Gln Leu Arg Glu Arg
                  20
 5
      (2) INFORMATION FOR SEQ ID NO: 761:
10
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 26 amino acids
                    (B) TYPE: amino acid
                  (D) TOPOLOGY: linear
            (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 761:
15
      Pro Ala Phe Gln Tyr Ser Ser His Val Ser Leu Gln Ala Ser Ser Gly
      His Met Trp Gly Thr Phe Arg Phe Glu Arg
20
                20
      (2) INFORMATION FOR SEQ ID NO: 762:
25 .
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 11 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
30
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 762:
      Ser Leu Cys Cys Pro Glu Gly Ala Glu Gly Cys
                        5
35
      (2) INFORMATION FOR SEQ ID NO: 763:
             (i) SEQUENCE CHARACTERISTICS:
40
                    (A) LENGTH: 12 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 763:
45
     Gln Leu Lys Lys Thr His Tyr Asp Arg Pro Cys Pro
       1
50
      (2) INFORMATION FOR SEQ ID NO: 764:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 12 amino acids
                    (B) TYPE: amino acid
55
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 764:
      Gln Leu Lys Lys Thr His Tyr Asp Arg Pro Cys Pro
60
```

	(2)	INF	ORMA	rion	FOR	SEQ	ID 1	10: 7	765:							
5			(i)	' (	ENCE A) L B) T D) T	engt Ype :	H: 1 ami	70 a no a	mino cid		ds				-	
10			(xi)	SEQ	-					EQ I	D NO	: 76	5:			
ıo	Ala 1	Gln	Arg	Lys	Lys 5	Glu	Met	Val	Leu	Ser 10	Glu	Lys	Val	Ser	Gln 15	Leu
15	Met	Glu	Trp	Thr 20	Asn	Lys	Arg	Pro	Val 25	Ile	Arg	Met	Asn	Gly 30	Asp	Lys
	Phe	Arg	Arg 35	Leu	Val	Lys	Ala	Pro 40	Pro	Arg	Asn	Tyr	Ser 45	Val	Ile	Val
20	Met	Phe 50	Thr	Ala	Leu	Gln	Leu 55	His	Arg	Gln	Cys	Val 60	Val	Cys	Lys	Gln
25	Ala 65	Asp	Glu	Glu	Phe	Gln -70	Ile	Leu	Ala	Asn	Ser 75	Trp	Arg	Tyr	Ser	Ser 80
	Ala	Phe	Thr	Asn	Arg 85	Ile	Phe	Phe	Ala	Met 90	Val	Asp	Phe	Asp	Glu 95	Gly.
30	Ser	Asp	Val	Phe 100	Gln	Met	Leu	Asn	Met 105	Asn	Ser	Ala	Pro	Thr 110	Phe	Ile
	Asn	Phe	Pro 115	Ala	Lys	Gly	Lys	Pro 120	Lys	Arg	Gly	Asp	Thr 125	Tyr	Glu	Leu
35	Gln	Val 130	Arg	Gly	Phe	Ser	Ala 135	Glu	Gln	Ile	Ala	Arg 140	Trp	Ile	Ala	Asp
40	Arg 145	Thr	Asp	Val	Asn	Ile 150	Arg	Val	Ile	Arg	Pro 155	Pro	Asn	Met	Ala	Ala 160
10	Arg	Trp	Arg	Phe	Trp 165	Cys	Val	Ser	Val	Thr 170						
45	(2)	INF	ORMA'	TION	FOR	SEQ	ID I	NO: '	766:							
50	(2) INFORMATION FOR SEQ ID NO: 766:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 15 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 766:								•							
55	Met 1	Val	Val	Ala	Leu 5	Leu	Ile	Val	Cys	Asp 10	Val	Pro	Ser	Ala	Ser 15	

(2) INFORMATION FOR SEQ ID NO: 767:

```
(i) SEQUENCE CHARACTERISTICS:
                   (A) LENGTH: 16 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
 5
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 767:
     Ala Gln Arg Lys Lys Glu Met Val Leu Ser Glu Lys Val Ser Gln Leu
            . 5
10
15
      (2) INFORMATION FOR SEQ ID NO: 768:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 17 amino acids
                    (B) TYPE: amino acid
20
                (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 768:
     Met Glu Trp Thr Asn Lys Arg Pro Val Ile Arg Met Asn Gly Asp Lys
                       5
       1
25
     Phe
30
      (2) INFORMATION FOR SEQ ID NO: 769:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 56 amino acids
35
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 769:
      Arg Arg Leu Val Lys Ala Pro Pro Arg Asn Tyr Ser Val Ile Val Met
40
              . 5
                                          10
      Phe Thr Ala Leu Gln Leu His Arg Gln Cys Val Val Cys Lys Gln Ala
                                     25
45
      Asp Glu Glu Phe Gln Ile Leu Ala Asn Ser Trp Arg Tyr Ser Ser Ala
      Phe Thr Asn Arg Ile Phe Phe Ala
50
      (2) INFORMATION FOR SEQ ID NO: 770:
55
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 31 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 770:
60
```

736

Met Val Asp Phe Asp Glu Gly Ser Asp Val Phe Gln Met Leu Asn Met

Asn Ser Ala Pro Thr Phe Ile Asn Phe Pro Ala Lys Gly Lys Pro

5

(2) INFORMATION FOR SEQ ID NO: 771: 10 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 37 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 771: 15 Lys Arg Gly Asp Thr Tyr Glu Leu Gln Val Arg Gly Phe Ser Ala Glu Gln Ile Ala Arg Trp Ile Ala Asp Arg Thr Asp Val Asn Ile Arg Val 20 25 Ile Arg Pro Pro Asn 35 25 (2) INFORMATION FOR SEQ ID NO: 772: 30 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 44 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 772: 35 Tyr Ala Gly Pro Leu Met Leu Gly Leu Leu Leu Ala Val Ile Gly Gly Leu Val Tyr Leu Arg Arg Val Ile Trp Asn Phe Ser Leu Ile Lys Leu 40 Asp Gly Leu Leu Gln Leu Cys Val Leu Cys Leu Leu 45 (2) INFORMATION FOR SEQ ID NO: 773: (i) SEQUENCE CHARACTERISTICS: 50 (A) LENGTH: 17 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 773: 55 Asp Ala Val Phe Lys Gly Phe Ser Asp Cys Leu Leu Lys Leu Gly Asp 5 10 Ser 60

	(2) INFORMATION FOR SEQ ID NO: 774:
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 20 amino acids
	<ul><li>(B) TYPE: amino acid</li><li>(D) TOPOLOGY: linear</li><li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 774:</li></ul>
10	Cys Gln Glu Gly Ala Lys Asp Met Trp Asp Lys Leu Arg Lys Glu Ser
15	Lys Asn Leu Asn 20
20	(2) INFORMATION FOR SEQ ID NO: 775:  (i) SEQUENCE CHARACTERISTICS:
	(A) LENGTH: 16 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear
25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 775:
	Val Leu Leu Val Ser Leu Ser Ala Ala Leu Ala Thr Trp Leu Ser Phe 1 5 10 15
30	
35	(2) INFORMATION FOR SEQ ID NO: 776:
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 48 amino acids
40	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 776:
45	Met Gly Leu Lys Leu Asn Gly Arg Tyr Ile Ser Leu Ile Leu Ala Val 1 5 10 15
43	Gln Ile Ala Tyr Leu Val Gln Ala Val Arg Ala Ala Gly Lys Cys Asp 20 25 30
50	Ala Val Phe Lys Gly Phe Ser Asp Cys Leu Leu Lys Leu Gly Asp Ser 35 40 45
55	
	(2) INFORMATION FOR SEQ ID NO: 777:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 90 amino acids

•	(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 777:															
5	Pro 1	Ala	Ala	Trp	Asp 5	Asp	Lys	Thr	Asn	Ile 10	Lys	Thr	Val	Cys	Thr 15	тут
10	Trp (	Glu	Asp	Phe 20	His	Ser	Cys	Thr	Val 25	Thr	Ala	Leu	Thr <sub>.</sub>	Asp 30	Cys	Gln
10	Glu (	Gly	Ala 35	Lys	Asp	Met	Trp	Asp 40	Lys	Leu	Arg	Lys	Glu 45	Ser	Lys	Asn
15	Leu i	Asn 50	Ile	Gln	Gly	Ser	Leu 55	Phe	Glu	Leu	Cys	Gly 60	Ser	Gly	Asn	Gly
	Ala 2 65	Ala	Gly	Ser	Leu	Leu 70	Pro	Ala	Phe	Pro	Val 75		Leu	Val	Ser	Leu 80
20	Ser 2	Ala	Ala	Leu	Ala 85	Thr	Trp	Leu	Ser	Phe 90					,	
25	(2)		ORMAT	SEQU	ENCE	СНА	RACT	ERIS	TICS	:	de					
30			(xi)	(	B) T	YPE:	ami : OGY	no a lin	cid ear			: 77	8:			
35	Met 1	Gly	Leu	Lys	Leu 5	Asn	Gly	Arg	Tyr	Ile 10	Ser	Leu	Ile	Leu	Ala 15	Val
	Gln	Ile	Ala	Tyr 20	Leu	Val	Gln	Ala	Val 25		Ala	Ala	Gly	Lys 30		Asp
40	Ala	Val	Phe 35		Gly	Phe	Ser	Asp 40		Leu	Leu	Lys	Leu 45		Asp	Ser
	Xaa	Xaa 50	Xaa	Xaa	Xaa	Pro	Ala 55		Trp	Asp	Asp	Lys 60		Asn	Ile	Lys
45	Thr 65	Val	Cys	Thr	Туг	Trp 70		Asp	Phe	His	Ser 75		Thr	Val	Thr	Ala 80
50	Leu	Thr	Asp	Cys	Gln 85		Gly	Ala	Lys	Asp 90		Trp	Asp	Lys	Leu 95	
50	Lys	Glu	Ser	Lys 100		Leu	Asn	ıle	Gln 105		Ser	Leu	Ph∈	Glu 110		Cy:
55	Gly	Ser	Gly 115		Gly	Ala	Ala	Gly 120		Leu	Leu	Pro	125		Pro	Va.

	(2) INFORMATION FOR SEQ ID NO: 779:
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 34 amino acids</li> <li>(B) TYPE: amino acid</li> <li>(D) TOPOLOGY: linear</li> <li>(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 779:</li> </ul>
10	Met Asn Ser Ala Ala Gly Phe Ser His Leu Asp Arg Arg Glu Arg Val  1 5 10 15
15	Leu Lys Leu Gly Glu Ser Phe Glu Lys Gln Pro Arg Cys Ala Ser Thr 20 25 30
	Leu Cys
20	(2) INFORMATION FOR SEQ ID NO: 780:
25	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 28 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 780:
30	Thr Ile Tyr Pro Thr Glu Glu Glu Leu Gln Ala Val Gln Lys Ile Val 1 5 10 15
	Ser Ile Thr Glu Arg Ala Leu Lys Leu Val Ser Asp 20 25
35	(2) INFORMATION FOR SEQ ID NO: 781:
40	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 30 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 781:
45	Arg Ala Leu Lys Gly Val Leu Arg Val Gly Val Leu Ala Lys Gly Leu 1 5 10 15
50	Leu Leu Arg Gly Asp Arg Asn Val Asn Leu Val Leu Leu Cys 20 25 30
	(2) INFORMATION FOR SEQ ID NO: 782:
55	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 39 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear
60	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 782:

PCT/US98/11422

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Ala Leu Ala Ala Leu Arg His Ala Lys Trp Phe Gln Ala Arg Ala Asn
     Gly Leu Gln Ser Cys Val Ile Ile Ile Arg Ile Leu Arg Asp Leu Cys
5
                                      25
     Gln Arg Val Pro Thr Trp Ser
             , 35
10
      (2) INFORMATION FOR SEQ ID NO: 783:
             (i) SEQUENCE CHARACTERISTICS:
15
                    (A) LENGTH: 17 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 783:
      Gly Asp Ala Leu Arg Arg Val Phe Glu Cys Ile Ser Ser Gly Ile Ile
20
                                      . 10
      Leu
25
      (2) INFORMATION FOR SEQ ID NO: 784:
             (i) SEQUENCE CHARACTERISTICS:
30
                     (A) LENGTH: 16 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 784:
35
      Leu Ala Phe Arg Gln Ile His Lys Val Leu Gly Met Asp Pro Leu Pro
                                           10
                        5
                               - ·
40
       (2) INFORMATION FOR SEQ ID NO: 785:
45
              (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 342 amino acids
                   . (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 785:
 50
      Thr. Ile Tyr Pro Thr Glu Glu Glu Leu Gln Ala Val Gln Lys Ile Val
                                           10
       Ser Ile Thr Glu Arg Ala Leu Lys Leu Val Ser Asp Ser Leu Ser Glu
 55
                    20
       His Glu Lys Asn Lys Asn Lys Glu Gly Asp Asp Lys Lys Glu Gly Gly
                                    40
 60
```

	Lys	Asp 50	Arg	Ala	Leu	Lys	Gly 55	Val	Leu	Arg	Val	Gly 60	Val	Leu	Ala	Lys
5	Gly 65	Leu	Leu	Leu	Arg	Gly 70	Asp	Arg	Asn	Val	Asn 75	Leu	Val	Leu	Leu	Cys 80
	Ser	Glu	Lys	Pro	Ser 85	Lys	Thr	Leu	Leu	Ser 90	Arg	Ile	Ala	Glu	Asn 95	Leu
10	Pro	Lys	Gln	Leu 100	Ala	Val	Ile	Ser	Pro 105	Glu	Lys	Tyr	Asp	Ile 110	Lys	Cys
15	Ala	Val	Ser 115	Glu	Ala	Ala	Ile	Ile 120	Leu	Asn	Ser	Суѕ	Val 125	Glu	Pro	Lys
	Met	Gln 130	Val	Thr	Ile	Thr	Leu 135	Thr	Ser	Pro	Ile	Ile 140	Arg	Glu	Glu	Asn
20	Met 145	Arg	Glu	Gly	Asp	Val 150	Thr	Ser	Gly	Met	Val 155	Lys	Asp	Pro	Pro	Asp 160
•					165			Leu		170					175	
25				180					185					190		Ile
30			195					200					205			Ser
		210	1				215	•				220				Ser
35	225					230					235					Phe 240
					245					250	)				255	
40	_			260	)				265	5				270		Asp
45			275	5				280	)		•		285	i		Leu
		290	)				295	5				300				Pro
50	309	5				310	)				319	5				320
	Ası	se:	r Ası	Gly	7 Val 325		Gl:	y Phe	e Glu	33		ı Gly	y Lys	. Lys	Asr 335	Lys 5
55	Ly:	s As <sub>l</sub>	р Ту:	r Ası 340		n Phe										

```
(i) SEQUENCE CHARACTERISTICS: .
                    (A) LENGTH: 24 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
5
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 786:
     Met Gly Ser Gln His Ser Ala Ala Ala Arg Pro Ser Ser Cys Arg Arg
                                           10
10
     Lys Gln Glu Asp Asp Arg Asp Gly
                  20
15
      (2) INFORMATION FOR SEQ ID NO: 787:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 30 amino acids
20
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 787:
      Leu Leu Ala Glu Arg Glu Gln Glu Glu Ala Ile Ala Gln Phe Pro Tyr
25
      Val Glu Phe Thr Gly Arg Asp Ser Ile Thr Cys Leu Thr Cys
30
      (2) INFORMATION FOR SEQ ID NO: 788:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 34 amino acids
35
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 788:
      Gln Gly Thr Gly Tyr Ile Pro Thr Glu Gln Val Asn Glu Leu Val Ala
40
                                         - 10
                        5 ·
      Leu Ile Pro His Ser Asp Gln Arg Leu Arg Pro Gln Arg Thr Lys Gln
45
      Tyr Val
50
       (2) INFORMATION FOR SEQ ID NO: 789:
              (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 55 amino acids
55
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 789:
      Ala Arg Leu Asn Val Gly Arg Glu Ser Leu Lys Arg Glu Met Leu Lys
 60
                         5
                                            10
```

```
Ser Gln Gly Val Lys Val Ser Glu Ser Pro Met Gly Ala Arg His Ser
                                      25
 5
      Ser Trp Pro Glu Gly Ala Ala Phe Cys Lys Lys Val Gln Gly Ala Gln
                                  40
      Met Gln Phe Pro Pro Arg Arg
           50
10
      (2) INFORMATION FOR SEQ ID NO: 790:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 15 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 790:
20
      Ala Arg Leu Asn Val Gly Arg Glu Ser Leu Lys Arg Glu Met Leu
                        5
25
      (2) INFORMATION FOR SEQ ID NO: 791:
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 20 amino acids
30
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 791:
      Leu Lys Ser Gln Gly Val Lys Val Ser Glu Ser Pro Met Gly Ala Arg
35
                                          10
             5
      His Ser Ser Trp
40
      (2) INFORMATION FOR SEQ ID NO: 792:
             (i) SEQUENCE CHARACTERISTICS:
45
                     (A) LENGTH: 17 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 792:
50
      Ala Phe Cys Lys Lys Val Gln Gly Ala Gln Met Gln Phe Pro Pro Arg
                                          10
      Arg
55
      (2) INFORMATION FOR SEQ ID NO: 793:
60
              (i) SEQUENCE CHARACTERISTICS:
```

```
(A) LENGTH: 17 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NC: 793:
5
     Ala Phe Cys Lys Lys Val Gln Gly Ala Gln Met Gln Phe Pro Pro Arg
                                           10
     Arg
10
      (2) INFORMATION FOR SEQ ID NO: 794:
15
             (i) SEQUENCE CHARACTERISTICS:
                    (A) LENGTH: 37 amino acids
                    (B) TYPE: amino acid
                    (D) TOPOLOGY: linear
             (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 794:
20
      Val Gln Val Leu Glu Gln Leu Thr Asn Asn Ala Val Ala Glu Ser Arg
       1
      Phe Asn Asp Ala Ala Tyr Tyr Trp Met Leu Ser Met Gln Cys Leu
25
                                       25
      Asp Ile Ala Gln Asp
               35
30
      (2) INFORMATION FOR SEQ ID NO: 795:
35
             (i) SEQUENCE CHARACTERISTICS:
                     (A) LENGTH: 34 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NC: 795:
40
      Pro Ala Gln Lys Asp Thr Met Leu Gly Lys Phe Tyr His Phe Gln Arg
      Leu Ala Glu Leu Tyr His Gly Tyr His Ala Ile His Arg His Thr Glu
45
                                        25
      Asp Pro
50
       (2) INFORMATION FOR SEQ ID NO: 796:
              (i) SEQUENCE CHARACTERISTICS:
55
                     (A) LENGTH: 27 amino acids
                     (B) TYPE: amino acid
                     (D) TOPOLOGY: linear
              (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 796:
       Leu Ala Lys Gln Ser Lys Ala Leu Gly Ala Tyr Arg Leu Ala Arg His
 60
```

	1.		5		10		•	15
5	Ala T	yr Asp Lys 20		Gly Leu	Tyr Ile 25	Pro		· .
	(2) I	NFORMATION	FOR SEQ	ID NO:	797:			
10		(i) SEQ	JENCE CHA (A) LENGT (B) TYPE: (D) TOPOL	H: 36 am amino a	ino acid cid	ls		,
15		(xi) SE	QUENCE DE	SCRIPTIO	N: SEQ I			Arg Ala
20	1 Lys F	ro Phe His	s Asp Ser	Glu Glu		Pro Leu	Cys Tyr 30	
	Ser 1	hr Asn Ası 35	1					•
25	(2) 1	NFORMATIO	n for seq	ID NO:	798:			
30			UENCE CHA (A) LENGT (B) TYPE: (D) TOPOI QUENCE DE	TH: 73 and and and and and and and and and and	mino ació acid near		8:	
35	Pro I	Leu Leu As	n Asn Leu 5	Gly Asn	Val Cys		.Cys Arg	Gln Pro 15
40		lle Phe Se 2 Leu Glu Gl	0	Thr Asp	25 Glu Glu		30 Ser Leu	
45	Leu (	35 Glu Val Le 50	u Arg Pro	40 Lys Arg · 55		Arg Gln 60		Ile Cys
 50	Lys (	Gln Gln Le	u Pro Asr 70		: Gly			
50	· (2) :	INFORMATIC	n for sec	ID NO:	799:			
55			(B) TYPE	TH: 29 ar : amino a LOGY: li	mino aci acid near		99:	
60	Met	Pro Tyr Al	a Gln Tri	o Leu Ala	a Glu Ası	n Asp Arg	Phe Glu	Glu Ala

	1		5		10			15
5	Gln Lys	s Ala Phe 1 20	His Lys A	la Gly A	rg Gln A 25	rg Glu	Ala	
	(2) IN	FORMATION :	FOR SEQ I	D NO: 80	00:			
10		(E	LENGTH: 3) TYPE: 6	: 36 ami amino ac GY: line	no acids id ar	NO+ 800	١.	
15		(XI) SEQU	ENCE DESC	RIFIION	. SEQ ID	140. 000		
	Phe Se	r Val His	Arg Pro G	Glu Thr	Leu Phe I	Asn Ile	Ser Arg	Phe Leu 15
20	Leu Hi	s Ser Leu 20	Pro Lys <i>P</i>	Asp Thr	Pro Ser ( 25	Sly Ile	Ser Lys 30	Val Lys
	Ile Le	u Phe Thr 35			÷	-		

A. The indications made below relate to the microorganism referron page 161 , line N/A							
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet						
Name of depositary institution  American Type Culture Collection							
Address of depositary institution (including postal code and count	n)						
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	•						
Date of deposit March 27, 1997	Accession Number 97979						
C. ADDITIONAL INDICATIONS (leave blank if not applicable)	le) This information is continued on an additional sheet						
D. DESIGNATED STATES FOR WHICH INDICATION	NS ARE MADE (if the indications are not for all designated States)						
·							
E. SEPARATE FURNISHING OF INDICATIONS (leave	blank if not applicable)						
	Bureau later (specify the general nature of the indications, e.g., "Accession						
For receiving Office use only	For International Bureau use only						
This sheet was received with the intermedianal application	This sheet was received by the International Bureau on:  Authorized officer						
0 4 JUN 1998							

A. The indications made below relate to the microorganism referred to in the description on page 162 . line N/A						
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet					
Name of depositary institution  American Type Culture Coll	ection					
Address of depositary institution (including postal code and country	(بر					
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America						
Date of deposit April 4, 1997	Accession Number 97974					
C. ADDITIONAL INDICATIONS (leave blank if not applicable	This information is continued on an additional sheet					
D. D. Colonia and						
D. DESIGNATED STATES FOR WHICH INDICATION	NS ARE MADE (if the indications are not for all designated States)					
·						
E. SEPARATE FURNISHING OF INDICATIONS (leave						
The indications listed below will be submitted to the International E Number of Deposit")	Bureau later (specify the general nature of the indications, e.g., "Accession					
	•					
·						
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Authorized officer	Authorized officer					
· · · · · · · · · · · · · · · · · · ·						

A. The indications made below relate to the microorganism referred to in the description on page 162 . line N/A						
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet					
Name of depositary institution  American Type Culture Col	lection					
Address of depositary institution (including postal code and country	(ער					
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America						
•						
Date of deposit May 29, 1997	Accession Number 209080					
C. ADDITIONAL INDICATIONS (leave blank if not applicable)	le) This information is continued on an additional sheet					
	•					
D. DESIGNATED STATES FOR WHICH INDICATION	NS ARE MADE (if the indications are not for all designated States)					
E SEDADATE DIDNISHING OF INDICATIONS						
E. SEPARATE FURNISHING OF INDICATIONS (leave The indications listed below will be submitted to the International)	blank if not applicable)  Bureau later (specify the general nature of the indications. e.g "Accession					
Number of Deposit")	topology in general mains of the maintains. e.g., Accession					
·						
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Authorized officer (1980) CCS-67 (7)	Authorized officer					

A. The indications made below relate to the microorganism refer on page 164 , line N/A	
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Co	Ilection
Address of depositary institution (including postal code and coun	ıry)
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
Date of deposit December 3, 1997	Accession Number 209511
C. ADDITIONAL INDICATIONS (leave blank if not applicate	ble) This information is continued on an additional sheet
•.	
D. DESIGNATED STATES FOR WHICH INDICATIO	NS ARE MADE (if the indications are not for all designated States)
	,
·	
E. SEPARATE FURNISHING OF INDICATIONS (leave	•
The indications listed below will be submitted to the International Number of Deposit")	Bureau later (specify the general nature of the indications, e.g., "Accession
	•
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Authorized officer (703) 205-3747	Authorized officer
0 4 JUN 1998	

A. The indications made below relate to the microorganism refer on page 167 . line N/A	
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution  American Type Culture Co	llection
Address of depositary institution (including postal code and count	ניזי)
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
Date of deposit April 4, 1997	Accession Number 97975
C. ADDITIONAL INDICATIONS (leave blank if not applicate	ble) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIO	NS ARE MADE (if the indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (leave	e blank if not applicable)
The indications listed below will be submitted to the International Number of Deposit')	Bureau later (specify the general nature of the indications, e.g., "Accession
	•
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This sheet was received with the international application  Authorized officer	This sheet was received by the International Bureau on:
0 4 JUN 1998	- Additionated Officer

A. The indications made below relate to the microorganism referred to in the description on page 167 , line N/A		
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	
Name of depositary institution  American Type Culture Collection		
Address of depositary institution (including postal code and country)		
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit May 29, 1997	Accession Number 209081	
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet		
	. •	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)		
	and the state of t	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)		
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")		
For receiving Office use only	Par International D	
This sheet was received with the international application	For International Bureau use only  This sheet was received by the International Bureau on:	
Authorized officer 1.0.5 1.0 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	Authorized officer	

A. The indications made below relate to the microorganism referred to in the description on page 171 . line N/A		
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	
Name of depositary institution  American Type Culture Col	llection	
Address of depositary institution (including postal code and count	ry)	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit April 4, 1997	Accession Number 97976	
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet		
D. DESIGNATED STATES FOR WHICH INDICATION	NS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)		
The indications listed below will be submitted to the International I Number of Deposit")	Bureau later (specify the general nature of the indications. e.g "Accession	
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This sheet was received with the international application cialist	This sheet was received by the International Bureau on:	
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A. The indications made below relate to the microorganism referred to in the description on page 172 . line N/A		
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	
Name of depositary institution American Type Culture Collection		
Address of depositary institution (including postal code and country)		
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit April 4, 1997	Accession Number 97977	
C. ADDITIONAL INDICATIONS (leave blank if not applicab	le) This information is continued on an additional sheet	
	·	
D. DESIGNATED STATES FOR WHICH INDICATION	NS ARE MADE (if the indications are not for all designated States)	
States)		
•		
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)		
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications as a "Accession"		
Number of Deposit")		
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Authorized officer	Authorized officer	
0 4 JUN 1998		

A. The indications made below relate to the microorganism referred to in the description on page 172 . line N/A			
IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet			
Name of depositary institution American Type Culture Co	ollection		
Address of depositary institution (including postal code and cour	niry)		
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America			
Date of deposit May 29, 1997	Accession Number 209082		
C. ADDITIONAL INDICATIONS (leave blank if not application)	able) This information is continued on an additional sheet		
D. DESIGNATED STATES FOR WHICH INDICATION	ONS ARE MADE (if the indications are not for all designated States)		
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A. The indications made below relate to the microorganism referred to in the description on page 176 , line N/A .		
DENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet		
Name of depositary institution  American Type Culture Collection		
Address of depositary institution (including postal code and country)		
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit April 28, 1997 Accession Number 209007		
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B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	
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10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
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A. The indications made below relate to the microorganism referred to in the description on page 179 , line N/A			
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A. The indications made below relate to the microorganism referred to in the description on page 182 , line N/A		
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Address of depositary institution (including postal code and country	y)	
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A. The indications made below relate to the microorganism referred to in the description on page 186 , line N/A .		
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Name of depositary institution American Type Culture Collection		
Address of depositary institution (including postal code and country)		
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Name of depositary institution  American Type Culture Collection			
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10801 University Boulevard Manassas, Virginia 20110-2209 United States of America			
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#### What Is Claimed Is:

- 1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:
- (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, having biological activity;
  - (f) a polynucleotide which is a variant of SEQ ID NO:X;
  - (g) a polynucleotide which is an allelic variant of SEQ ID NO:X;
  - (h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;
- (i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.
- 2. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.
- 3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

- 4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.
- 5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.
- 6. The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.
- 7. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.
- 8. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.
  - 9. A recombinant host cell produced by the method of claim 8.
  - 10. The recombinant host cell of claim 9 comprising vector sequences.
- 11. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:
- (a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z, having biological activity;
- (c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (e) a secreted form of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (f) a full length protein of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

- (g) a variant of SEQ ID NO:Y;
- (h) an allelic variant of SEQ ID NO:Y; or
- (i) a species homologue of the SEQ ID NO:Y.
- 12. The isolated polypeptide of claim 11, wherein the secreted form or the full length protein comprises sequential amino acid deletions from either the C-terminus or the N-terminus.
- 13. An isolated antibody that binds specifically to the isolated polypeptide of claim 11.
- 14. A recombinant host cell that expresses the isolated polypeptide of claim 11.
  - 15. A method of making an isolated polypeptide comprising:
- (a) culturing the recombinant host cell of claim 14 under conditions such that said polypeptide is expressed; and
  - (b) recovering said polypeptide.
  - 16. The polypeptide produced by claim 15.
- 17. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 11 or the polypucleotide of claim 1.
- 18. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
- (a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation.
- 19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
- (a) determining the presence or amount of expression of the polypeptide of claim 11 in a biological sample; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

- 20. A method for identifying a binding partner to the polypeptide of claim 11 comprising:
  - (a) contacting the polypeptide of claim 11 with a binding partner; and
- (b) determining whether the binding partner effects an activity of the polypeptide.
  - 21. The gene corresponding to the cDNA sequence of SEQ ID NO:Y.
- 22. A method of identifying an activity in a biological assay, wherein the method comprises:
  - (a) expressing SEQ ID NO:X in a cell;
  - (b) isolating the supernatant;
  - (c) detecting an activity in a biological assay; and
  - (d) identifying the protein in the supernatant having the activity.
  - 23. The product produced by the method of claim 22.

#### PATENT COOPERATION TREATY

# **PCT**

# DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT (PCT Article 17(2)(a) and Rule 39)

Applicant's or agent's file reference PZ007PCT	IMPORTANT DECLARATI	ON Date of mailing (day/month/year)	
International application No.	International filing date (day/mont	h/year) (Earliest) Priority Date (day/month/year)	
PCT/US98/11422	04 JUNE 1998	06 JUNE 1997	
International Patent Classification (IPC) Please See Continuation Sheet.	or both national classification and II	oc 1	
Applicant HUMAN GENOME SCIENCES, INC			
This International Searching Authority hereby declares, according to Article 17(2)(a), that no international search report will be established on the international application for the reasons indicated below.			
1. The subject matter of the interest as Scientific theories.	emational application relates to:		
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c. plant varieties.	<del></del>		
d. animal varieties.		· .	
e. essentially biologica and the products of	l processes for the production of plant such processes.	s and animals, other than microbiological processes	
f. schemes, rules or m	ethods of doing business.		
g. schemes, rules or m	ethods of performing purely mental	acts.	
h. schemes, rules or m	ethods of playing games.	•	
i. methods for treatme	nt of the human body by surgery or	therapy.	
j. methods for treatme	nt of the animal body by surgery or	therapy.	
k. diagnostic methods	practiced on the human or animal bo	ody.	
1. mere presentations	of information.		
m. computer programs	for which this International Searchin	g Authority is not equipped to search prior art.	
2. The failure of the following meaningful search from being	parts of the international application g carried out:	to comply with prescribed requirements prevents a	
the description	the claims	the drawings	
3. X  The failure of the nucleotide and/or amino acid sequence listing to comply with the prescribed requirements prevents a meaningful search from being carried out:			
it does not comply with the prescribed standard			
it is not in the prescribed machine readable form			
4. Further comments: Please See Continuation Sheet.			
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Name and mailing address of the ISA/US  Authorized offices			
Name and mailing address of the ISA/US  Commissioner of Patents and Trademarks Box PCT  Washington, D.C. 20231  Authorized offices  BRIAN R. STANTON			
Facsimile No. (703) 305-3230	Telephone	No. (703) 308-0196	

Form PCT/ISA/203 (July 1992)\*

# DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT

International application No. PCT/US98/11422

The International Patent Classification (IPC) or National Classification and IPC are as listed below:

IPC(6): A01N 37/18, 43/04; C12Q 1/00, 1/02, 1/68; C12N 5/00, 5/06, 15/00, 15/06, 15/09, 15/10, 15/11; G01N 33/53

US CL.: 435, 4, 7.1, 69.1, 70.1, 71.1, 172.3, 243, 320.1, 325, 410; 514/2, 44; 530/350, 387.1

#### 4. Further Comments (Continued):

Applicant has not responded to the invitation to pay additional fees mailed on 04 August 1998. Therefore, the search would be conducted on the first appearing invention white includes claims 1-10, 14, and 15 in so far as these claims are drawn to the first ten (10) appearing nucleotide sequences. However, no meaningful search could be carried out on these sequences because the CRF that was received for this case on 15 June 1998 was technically defective and could not be used to conduct a search of the prior art.